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INTERIM REPORT February 25, 1993

FOR

BIOVENTING FIELD INITIATIVE

AT

HANSCOM AIR FORCE BASE, MASSACHUSETTS

to

Captain Catherine M. Vogel
Department of the Air Force
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INTERIM REPORT

BIOVENTING FIELD INITIATIVE

HANSCOM AIR FORCE BASE, MASSACHUSETTS

1.0 INTRODUCTION

This report describes the activities conducted at Hanscom Air Force Base (AFB), Massachusetts, as part of the Bioventing Field Initiative for the U.S. Air Force Center for Environmental Excellence (AFCEE) and the Environmental Quality Directorate of the Air Force Armstrong Laboratory. This report summarizes the results from the first phase of the study at Hanscom AFB. First-phase activities include a soil gas survey, air permeability test, in situ respiration tests, and installation of bioventing systems. The specific objectives of this Bioventing Field Initiative are described in the following section. Each site at the base is discussed individually, followed by a description of site activities at the background area.

1.1 Objectives

The purpose of this Bioventing Field Initiative is to measure the soil gas permeability and microbial activity at a contaminated site in order to evaluate the potential application of bioventing technology to remediate the site. The specific test objectives are stated below.

- A small-scale soil gas survey will be conducted to identify an appropriate location for installation of the bioventing system. Soil gas from the candidate site should exhibit high total petroleum hydrocarbon (TPH) concentrations, relatively low oxygen concentrations, and relatively high carbon dioxide concentrations. An uncontaminated background location also will be identified.
- The soil gas permeability of the soil and the air vent (well) radius of influence will be determined. To measure these parameters, air will be withdrawn or injected for approximately 8 hours at vent wells located in contaminated soils. Pressure changes will be monitored in an array of monitoring points.
- Immediately following the soil gas permeability test, an in situ respiration test will be conducted. Air will be injected into selected monitoring points to

aerate the soils. The in situ oxygen utilization and carbon dioxide production rates will be measured.

The data from the soil gas permeability and in situ respiration tests will be used to determine an air injection/withdrawal rate for the bioventing test. A blower will be selected, installed, and operated for 6 to 12 months, and periodic measurements of the soil gas composition will be made to evaluate the long-term effectiveness of bioventing.

1.2 Site Descriptions

Hanscom AFB is located in Bedford, Massachusetts. A schematic diagram of the base is shown in Figure 1. The dashed lines on the map indicate the direction from the main gate to each test site. Site H1 and Site H2 refer to Building 1639 and Building 1812, respectively. Groundwater at Hanscom AFB generally is encountered at 3 to 8 feet. The sites chosen for the bioventing test initiative are Building 1639 and Building 1812. A brief description of the sites is provided in Sections 1.2.1 and 1.2.2. A detailed description of the test sites is provided in the Test Plan in Appendix A.

1.2.1 Building 1639 Site

Building 1639 is the base fuel service station (Figure 2). There have been three reported petroleum releases (gasoline, #2 fuel oil, and waste lubricating oil) at this facility since November 1990. Soil contamination at Building 1639 ranges from 599 ppm to 4,400 ppm TPH.

1.2.2 Building 1812 Site

The site at Building 1812 was the location of a heating oil fuel tank (Figure 3). Leakage occurred from the tank, resulting in soil contamination at this site. The tank was replaced recently, and soil TPH concentrations as high as 2,830 mg/kg have been reported.

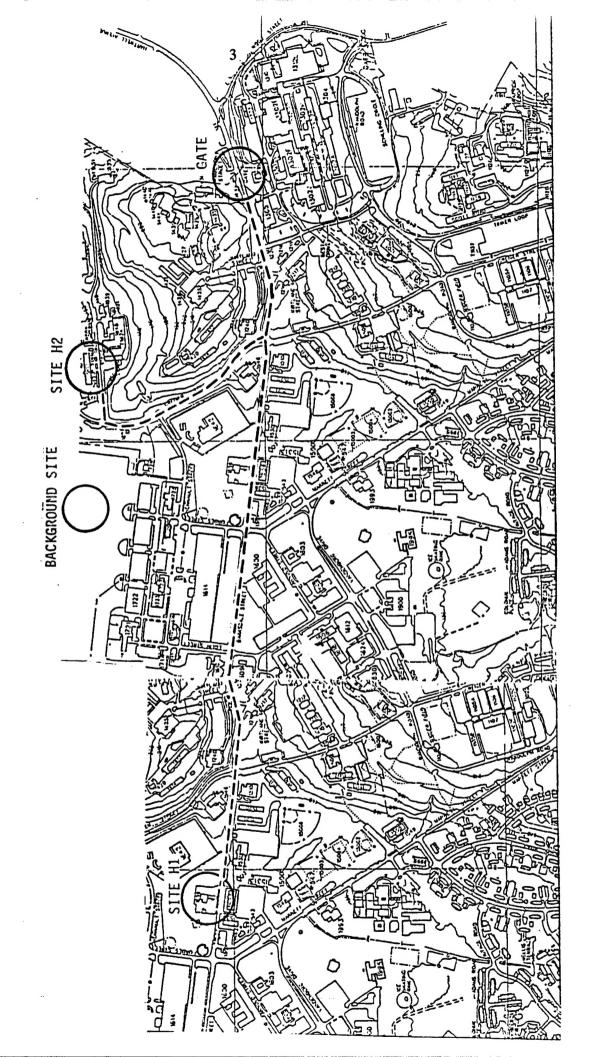


Figure 1. Schematic Diagram of Hanscom AFB

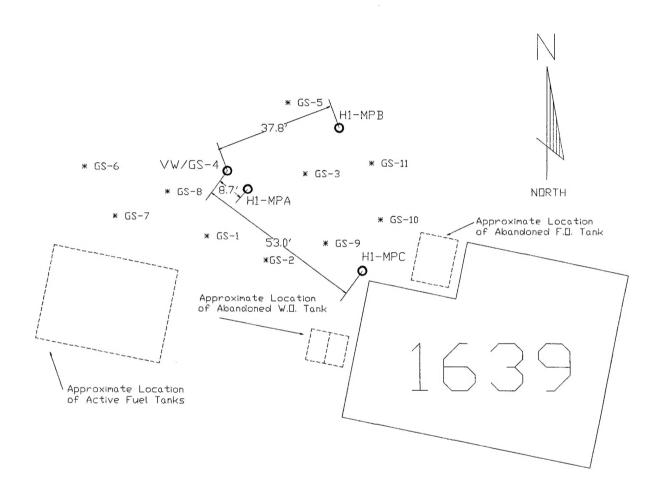


Figure 2. Schematic Diagram of Building 1639 at Hanscom AFB (GS - Soil Gas Survey Point; MP - Monitoring Point)

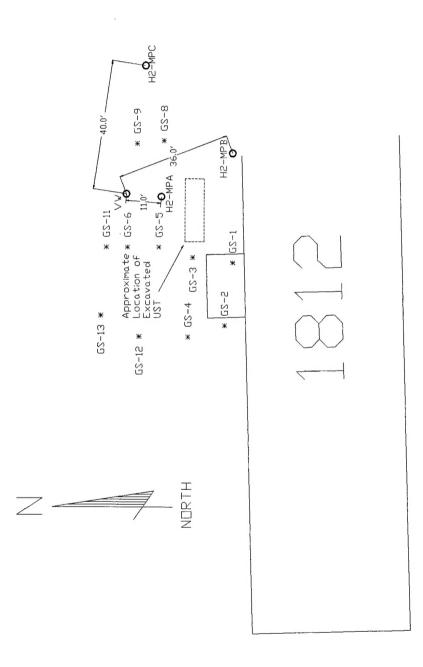


Figure 3. Schematic Diagram of Building 1812 at Hanscom AFB (GS - Soil Gas Survey Point; MP - Monitoring Point)

2.0 BUILDING 1639

2.1 Chronology of Events and Site Activities

2.1.1 Groundwater Measurements

One groundwater monitoring well (12BO3) was present at Building 1639. Groundwater level was measured at this well on September 30, 1992 and was recorded at 5.75 feet.

2.1.2 Soil Gas Survey

A suitable site for the bioventing demonstration should have soil gas characteristics of high TPH, low oxygen, and high carbon dioxide concentrations. This composition of soil gas would indicate that oxygen-limiting conditions for microbial activity are present and that the introduction of air may enhance biodegradation of TPH.

On September 30, 1992, a limited soil gas survey was conducted at Building 1639. Soil gases were sampled by driving a %-inch-diameter stainless steel probe into the soil with a hammer drill. Soil gas was withdrawn with a vacuum pump and analyzed for oxygen, carbon dioxide, and TPH.

Measurements of oxygen and carbon dioxide in the soil gas were made with a GasTech Model 32520X with oxygen and carbon dioxide ranges of 0 to 25%. The analyzer was calibrated daily against atmospheric oxygen, atmospheric carbon dioxide, a 10% oxygen calibration standard, and a 5% carbon dioxide calibration standard. TPH was measured with a GasTech Trace Techtor with TPH ranges from 0 to 100, 0 to 1,000, and 0 to 10,000 ppm. The GasTech Trace Techtor was calibrated daily against a 4,200 ppm hexane standard.

The soil gas probes were driven to depths ranging from 2.5 to 5.0 feet at several locations at Building 1639. Table 1 provides the initial concentrations of oxygen, carbon dioxide, and TPH for the various locations at Building 1639. Oxygen concentrations varied from 0.8 to 21%, whereas TPH concentrations ranged from 410 ppm to greater than 20,000 ppm. The oxygen concentrations in the soil gas indicate that some areas at this site are oxygen-limited and may respond to bioventing.

Table 1. Initial Soil Gas Composition at Building 1639

Soil Gas Survey (GS) Point	Depth (ft)	Oxygen (%)	Carbon Dioxide (%)	TPH (ppm)
GS-1	2.5	19.0	2.5	15,600
	5.0	20.2	0.70	6,000
GS-2	2.5	16.0¹	5.8	>20,000
	5.0	2.0	18.5	>20,000
GS-3	2.5	1.0	18.5	>20,000
	5.0	3.0	17.0	>20,000
GS-4	2.5	21.0¹	0.10	920
	5.0	1.0	23.0	>20,000
GS-5	2.5	16.2	5.8	660
	5.0	15.0	7.0	720
GS-6	2.5	20.0	1.8	410
GS-7	2.5	21.0	0.50	900
GS-8	2.5	19.5¹	1.0	5,600
	5.0	19.2¹	1.8	8,800
GS-9	2.5	4.0 ⁱ	13.5	>20,000
	5.0	0.801	19.5	>20,000
GS-10	2.5	13.5¹	6.8	>20,000
	5.0	0.801	17.0	>20,000
GS-11	2.5	16.0¹	4.3	>20,000

Pressure reading on sampling pump was high. Measured oxygen concentration may not be representative of actual soil gas oxygen concentrations. Actual oxygen concentration is likely to be lower.

2.1.3 Vent Well, Monitoring Point, and Thermocouple Installation

On October 3, 1992, one vent well (VW) and three monitoring points (MPs) were installed, and soil samples were collected for analyses. The monitoring points were labeled as follows: H1-MPA; H1-MPB; and H1-MPC. The locations of the vent well and monitoring points are shown in Figure 2. A cross section of the vent well and monitoring points showing site lithology and construction detail is shown in Figure 4.

The vent well was installed at a depth of 7.0 feet into an 8-inch-diameter borehole. The vent well consisted of Schedule 40 2-inch-diameter polyvinyl chloride (PVC) piping with 3.0 feet of tenslot screen. The annular space corresponding to the screened area of the well was filled with silica sand, and the annular space above the screened interval was filled with bentonite to prevent short-circuiting of air to or from the surface.

Soil gas probes consisted of ¼-inch tubing with a 1-inch-diameter, 6-inch screened area. The annular space corresponding to the screened area was filled with silica sand, whereas the interval between the screened areas was filled with bentonite, as was the annular space from the shallowest monitoring point to the ground surface. The monitoring points were installed at depths as follows:

- Monitoring point H1-MPA was installed at a depth of 5.0 feet into an 8-inch-diameter borehole. The monitoring point was screened to two depths: 2.5 and 5.0 feet.
- Monitoring point H1-MPB was installed at a depth of 5.0 feet into an 8-inch-diameter borehole. The monitoring point was screened to two depths: 2.5 and 5.0 feet.
- Monitoring point H1-MPC was installed at a depth of 6.0 feet into an 8-inch-diameter borehole. The monitoring point was screened to two depths: 3.5 and 6.0 feet.

A Type K thermocouple was installed with monitoring points H1-MPA-2.5' and H1-MPA-5.0'.

MPC

MPB

MPA

Vent Well

Figure 4. Cross Section of Vent Well and Monitoring Points at Building 1639 Showing Site Lithology and Construction Detail (not to scale)

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2.1.4 Soil and Soil Gas Sampling and Analyses

Split-spoon soil samples were collected at depths of 4.0 to 4.5 feet and 4.5 to 5.0 feet from the vent well borehole and were labeled H1-VW-4'-4.5' and H1-VW-4.5'-5.0', respectively. A soil sample also was collected from monitoring point A at a depth of 3.0 to 4.0 feet and was labeled H1-A-3'-4'. The samples were sent under chain of custody to Engineering-Science Inc., Berkeley Laboratory for analysis of benzene, toluene, ethylbenzene, and xylenes (BTEX); TPH; alkalinity; moisture content; pH; iron; total phosphorous; total Kjeldahl nitrogen; and particle size analysis.

Soil vapor samples were collected from all monitoring points and were labeled H1-A-2.5, H1-A-5, H1-B-2.5, H1-B-5, H1-C-3.5, and H1-C-6. These samples were sent under chain of custody to Air Toxics, Ltd., in Rancho Cordova, California, for analysis of BTEX and TPH.

2.1.5 Soil Gas Permeability and Radius of Influence

A detailed description of the method for conducting a soil gas permeability test, including equations to compute k, the soil gas permeability, is given in the Test Plan and Technical Protocol (Hinchee et al., 1992).

Prior to air injection, the monitoring points were allowed to set up for 24 hours. Air was injected with a portable 1-horsepower (HP) explosion-proof positive displacement blower unit. After air injection was initiated, pressure readings were taken approximately every 1 to 2 minutes for the first hour, then approximately every 10 minutes for the following hour. The HyperventilateTM computer model was used to calculate the soil gas permeability.

2.1.6 In Situ Respiration Test

Immediately following the soil gas permeability test, air containing approximately 1% helium was injected into the soil for approximately 24 hours, beginning on October 6. Air was injected concurrently into the background monitoring well to measure the natural biodegradation of organic material in the soil. The setup for the in situ respiration test is described in the Test Plan and Technical Protocol (Hinchee et al., 1992). The pump used for air injection was a 1/3-HP diaphragm pump. Air and helium were injected through the following monitoring points at the depths indicated: H1-MPA-5.0'; H1-MPB-2.5'; H1-MPB-5.0'; and H1-MPC-6.0'. After the air/helium injection was

turned off, the respiration gases were monitored periodically. The respiration test was terminated on October 10.

Helium concentrations were measured during the in situ respiration test to quantify helium leakage to or from the surface around the monitoring points. Helium loss over time is attributed to either diffusion or leakage. A rapid drop in helium concentration followed by a leveling is an indication of leakage. A gradual loss along with an apparent first-order curve is an indicator of diffusion. As a rough estimate, the diffusion of gas molecules is inversely proportional to the square root of the molecular weight of the gas. Based on molecular weights of 4 for helium and 32 for oxygen, helium gas diffuses about 2.8 times faster than oxygen, or the diffusion of oxygen is 0.35 times the rate of helium diffusion. As a general rule, we have found that if helium concentrations are at least 50 to 60% of the initial levels at test completion, measured oxygen uptake rates are representative. Greater helium loss indicates a problem, and oxygen utilization rates are not considered representative.

To compare data from one site to another, a stoichiometric relationship of the oxidation of the hydrocarbon was assumed. Hexane was used as the representative hydrocarbon for the organic contaminant. The stoichiometric relationship is given by:

$$C_6H_{14} + 9.5O_2 - 6CO_2 + 7H_2O$$
 (1)

Based on the utilization rates (% per day), the biodegradation rates in terms of milligrams as a hexane equivalent per kilogram of soil per day were computed using the equation below by assuming a soil porosity of 0.2 and a bulk density of 1,440 kg/m³.

$$K_{\beta} = \frac{-K_o A D_o C}{100}$$
 (2)

where: K_{g} = biodegradation rate (mg/kg/day)

K_o = oxygen utilization rate (percent per day)

A = volume of air/kg of soil, in this case 300/1,440 = 0.21

 D_o = density of oxygen gas (mg/L) assumed to be 1,330 mg/L

C = mass ratio of hydrocarbon to oxygen required for mineralization, assumed to be 1/3.5 from the above stoichiometric equation.

2.2 Results and Discussion

2.2.1 Soil and Soil Gas Analyses

Results of the soil analyses for BTEX and TPH at Building 1639 are presented in Table 2. The analytical report for this site is presented in Appendix B. Concentrations of the BTEX compounds in soil samples ranged from 0.015 mg/kg (ethylbenzene) up to 12 mg/kg (total xylenes), whereas TPH concentrations ranged from below the detection limit (<0.0040 mg/kg) to 22 mg/kg. The soil vapor analyses also showed similar measurements of BTEX and TPH, with concentrations of TPH ranging from 280 ppmv to 19,000 ppmv and from 0.11 ppmv (ethylbenzene) up to 67 ppmv (total xylenes) of the BTEX compounds (Table 2). The results of the soil chemistry analyses are summarized in Table 3.

2.2.2 Soil Gas Permeability and Radius of Influence

The raw data for the soil gas permeability test at Building 1639 are presented in Appendix C. Using the HyperventilateTM computer model, soil gas permeabilities were calculated at each of the monitoring points. These data are presented in Table 4. The soil gas permeability varied considerably, with values ranging from 24 darcy up to 5.4 x 10⁸ darcy. Typically, the radius of influence is calculated by plotting the log of the pressure change at a specific monitoring point versus the distance from the vent well. The radius of influence would then be the distance where 1 inch of water pressure can be measured. However, in this instance, 1 inch of water pressure was not achieved at any monitoring point (Figure 5); therefore, a radius of influence based on these specifications cannot be definitively determined at this site, other than to say it is less than 8.7 feet.

2.2.3 In Situ Respiration Test

The results of the in situ respiration test for Building 1639 are presented in Appendix D. Each figure in Appendix D illustrates the oxygen, carbon dioxide, and helium concentrations as a

Table 2. Results From Soil and Soil Gas Analyses for BTEX and TPH at Building 1639

Matrix	Sample Name	Benzene (mg/kg)	Toluene (mg/kg)	Ethylbenzene (mg/kg)	Total Xylenes (mg/kg)	TPH¹ (mg/kg)
Soil	H1-VW-4'-4.5'	0.048	0.020	0.015	0.020	22
	H1-VW-4.5'-5.0'	0.67	0.27	0.43	0.45	15
	H1-A-3'-4'	1.0	4.3	1.3	12	< 0.0040
Matrix	Sample Name	Benzene (ppmv)	Toluene (ppmv)	Ethylbenzene (ppmv)	Total Xylenes (ppmv)	TPH² (ppmv)
Soil Gas	H1-A-2.5	5.2	4.2	1.1	2.9	5,600
	H1-A-5	27	35	10	30	19,000
	H1-B-2.5	2.8	1.3	0.84	1.9	2,700
	H1-B-5	2.4	0.84	0.42	1.5	3,200
	H1-C-3.5	0.44	0.13	0.11	0.37	280
	H1-C-6	11	20	9.3	67	11,000

Referenced to a reference oil composed of a mixture of 2,2,4-trimethylpentane, *n*-hexadecane, and chlorobenzene.

² TPH referenced to jet fuel (molecular weight = 156).

Table 3. Results From Soil Chemistry Analyses at Building 1639

	Sample Name					
Parameter	H1-VW-4'-4.5'	H1-VW-4.5′-5.0′	H1-A-3'-4'			
Alkalinity (mg/kg CaCO ₃)	< 50	< 50	< 50			
Moisture (% by weight)	22.2	21.8	5.8			
рН	5.8	6.0	6.1			
Iron (mg/kg)	7,980	6,260	8,630			
Total Phosphorous (mg/kg)	370	290	300			
Total Kjeldahl Nitrogen (mg/kg)	1,100	730	70			
Particle Size Analysis (%)	Gravel: 11.5	Gravel: 0.5	Gravel: 26			
	Sand: 59	Sand: 72	Sand: 55			
	Silt: 27	Silt: 23	Silt: 16			
	Clay: 6	Clay: 4.5	Clay: 3			

Table 4. Results of Hyperventilate™ Soil Gas Permeability Analysis at Building 1639

Monitoring Point	Depth (ft)	Soil Gas Permeability (darcy)
H1-MPA	2.5	24
	5.0	130
Н1-МРВ	2.5	1,400
	5.0	6,200
H1-MPC	3.5	5.4 x 10 ⁸
	6.0	28

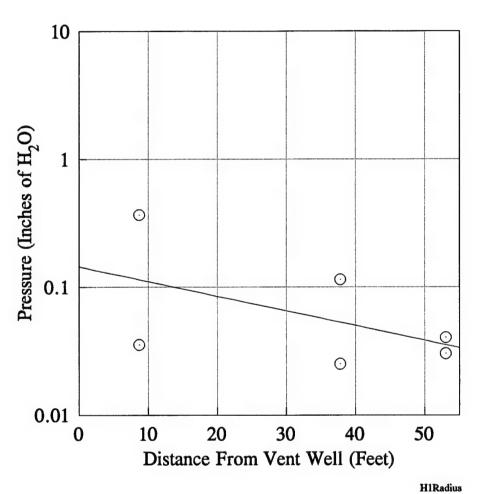


Figure 5. Radius of Influence at Building 1639

function of time. An example of typical oxygen utilization at this site is shown in Figure 6, where oxygen utilization and carbon dioxide production at monitoring point H1-MPA-5.0' is illustrated. A summary of the oxygen utilization and carbon dioxide production rates and corresponding biodegradation rates is shown in Table 5. The biodegradation rates measured at this site were fairly high, with rates ranging from 8.0 mg/kg/day to 27 mg/kg/day based on oxygen utilization, and from 0.48 mg/kg/day to 4.3 mg/kg/day based on carbon dioxide production.

Loss of helium was insignificant at all monitoring points, indicating that the monitoring points were well sealed and that the oxygen depletion observed was a result of biodegradation.

Soil temperatures were measured during the in situ respiration test. Temperatures during the test ranged from 17.9 to 19.3°C at monitoring point H1-MPA-2.5′ and from 18.7 to 20.7°C at monitoring point H1-MPA-5.0′.

2.2.4 Bioventing Demonstration

The decision was made to install a bioventing system at Building 1639. A 1-HP blower was installed at the site on October 14, 1992. Air injection was initiated on October 14 at a flowrate of 2.5 scfm.

Table 5. Oxygen Utilization and Carbon Dioxide Production Rates During the In Situ Respiration Test at Building 1639

Monitoring Point	Oxygen Utilization Rate (%/hour)	Biodegradation Rate (mg/kg/day)	Carbon Dioxide Production Rate (%/hour)	Biodegradation Rate (mg/kg/day)
Background	0.0073	0.14	0.0087	0.19
H1-MPA-5.0'	1.4	27	0.076	1.6
H1-MPB-2.5'	0.42	8.0	0.034	0.74
H1-MPB-5.0'	0.58	11	0.022	0.48
H1-MPC-6.0'	0.69	13	0.20	4.3

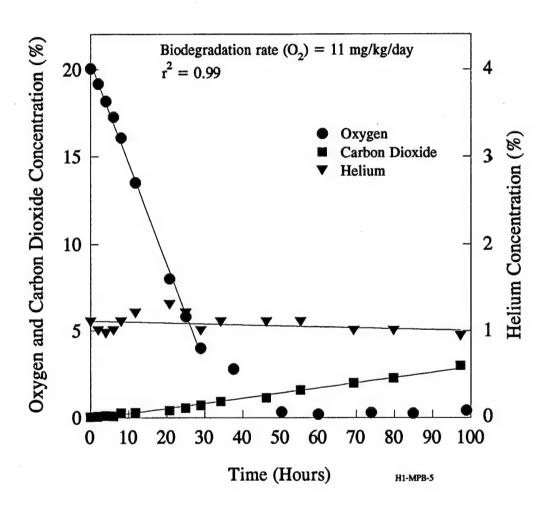


Figure 6. Oxygen Utilization and Carbon Dioxide Production During the In Situ Respiration Test at Monitoring Point H1-MPB-5.0'

3.0 BUILDING 1812

3.1 Chronology of Events and Site Activities

Although oxygen levels measured during the soil gas survey indicated that some areas at the Building 1812 site may be limited, oxygen concentrations measured at the permanent monitoring points were fairly high. Therefore, an in situ respiration test was not conducted at this site. However, because measurements taken during the soil gas survey indicated that some areas were oxygen-limited, a bioventing system was installed at the site in order to treat those areas. Other activities were conducted at the site according to the Test Plan and Technical Protocol (Hinchee et al., 1992).

3.1.1 Groundwater Measurements

Groundwater depth was measured at one monitoring well (HB-02) at Building 1812. The groundwater level was measured on October 1, 1992 and was recorded at 3.98 feet.

3.1.2 Soil Gas Survey

On October 1, 1992, a limited soil gas survey was conducted to locate a suitable test area at Building 1812. Soil gases were sampled by driving a %-inch-diameter stainless steel probe into the soil with a hammer drill. Soil gas was withdrawn with a vacuum pump and analyzed for oxygen, carbon dioxide, and TPH. Soil gas measurements were taken as described in Section 2.1.2.

The soil gas probes were driven to depths ranging from 2.0 to 5.0 feet at several locations at Building 1812. Table 6 provides the initial concentrations of oxygen, carbon dioxide, and TPH for the various locations at Building 1812. Relatively high concentrations of oxygen were found at most of the soil gas probes, with concentrations ranging from 3 to 20%. Relatively low concentrations of carbon dioxide (0.05 to 8.5%) and TPH (10 ppm to 800 ppm) were encountered. The oxygen concentrations in the soil gas indicate that some areas at this site are oxygen-limited and may respond to bioventing.

Table 6. Initial Soil Gas Composition at Building 1812

Monitoring Point	Depth (ft)	Oxygen (%)	Carbon Dioxide (%)	TPH (ppm)
GS-1	2.5	19.5	0.7	120
GS-2	2.5	20	0.05	10
	5.0	NS	NS	NS
GS-3	2.25	20.2	0.60	87
GS-4	2.5	17	0.30	40
	4.0	19.8¹	0.08	20
GS-5	2.5	15	3.8	220
	5.0	13	5.1	270
GS-6	2.5	16.5	1.3	120
	5.0	NS	NS	NS
GS-7	2.0	9.1	6.0	440
	3.5 - 4.0	7.0	7.2	480
GS-8	2.5	17.5	1.2	250
GS-9	2.0	17	1.4	280
GS-10	2.0	3.0	8.5	800
GS-11	2.0	18.8	0.40	180
	3.0	18.3	0.60	96
	4.0	18.9¹	2.7	220
GS-12	2.0	17	0.10	41
	3.0	18	0.70	120
	4.0	17.5	0.10	50

NS Not sampled. Groundwater was encountered at this depth.

Pressure reading on sampling pump was high. Measured oxygen concentration may not be representative of actual soil gas oxygen concentrations. The actual oxygen concentration is likely to be lower.

3.1.3 Vent Well, Monitoring Point, and Thermocouple Installation

On October 4, 1992, one vent well and three monitoring points were installed, and soil samples were collected for analyses. The monitoring points were labeled H2-MPA, H2-MPB, and H2-MPC. The locations of the vent well and monitoring points are shown in Figure 3. A cross section of the vent well and monitoring points showing site lithology and construction detail is shown in Figure 7.

The vent well was installed at a depth of 7.0 feet into an 8-inch-diameter borehole. The vent well consisted of Schedule 40 2-inch-diameter PVC piping with 3.6 feet of ten-slot screen. The annular space corresponding to the screened area of the well was filled with silica sand, whereas the annular space above the screened interval was filled with bentonite to prevent short-circuiting of air to or from the surface.

Soil gas probes consisted of ¼-inch tubing with a 1-inch-diameter, 6-inch screened area. The annular space corresponding to the screened area was filled with silica sand, whereas the interval between the screened areas was filled with bentonite, as was the annular space from the shallowest monitoring point to the ground surface. The monitoring points were installed at depths as follows:

- Monitoring point H2-MPA was installed at a depth of 5.0 feet into an 8-inch-diameter borehole. The monitoring point was screened to two depths: 2.5 and 5.0 feet.
- Monitoring point H2-MPB was installed at a depth of 7.0 feet into an 8-inch-diameter borehole. The monitoring point was screened to three depths: 2.5, 5.0, and 7.0 feet.
- Monitoring point H2-MPC was installed at a depth of 6.0 feet into an 8-inch-diameter borehole. The monitoring point was screened to three depths: 2.5, 4.5, and 6.0 feet.

A Type K thermocouple was installed with monitoring points H2-MPA-2.5' and H2-MPA-5.0'.

MPC

MPB

MPA

Vent Well

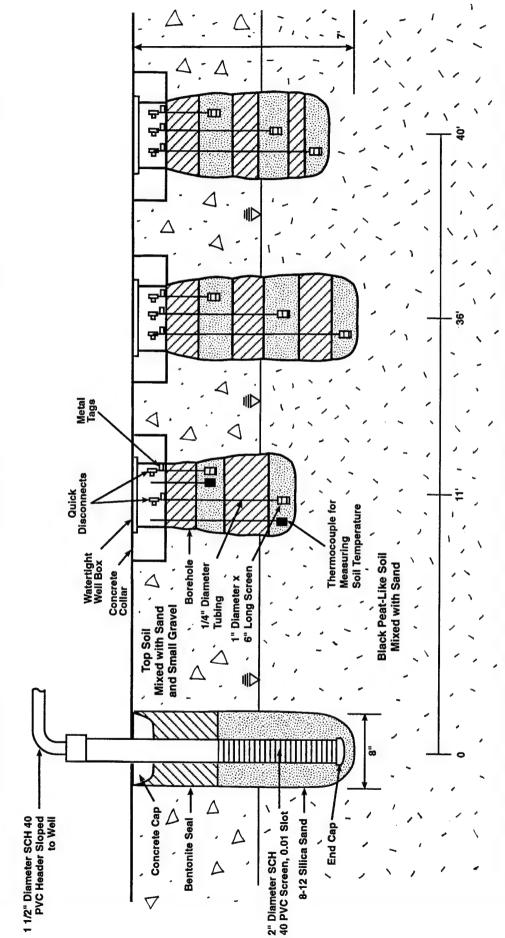


Figure 7. Cross Section of Vent Well and Monitoring Points at Building 1812 Showing Site Lithology and Construction Detail (not to scale)

F/Kittel11/h-2

3.1.4 Soil and Soil Gas Sampling and Analyses

Split-spoon soil samples were collected at depths of 3.0 to 3.5 feet and 4.0 to 4.5 feet from the vent well borehole and were labeled H2-VW-3'-3.5' and H2-VW-4'-4.5', respectively. The samples were sent under chain of custody to Engineering-Science Inc., Berkeley Laboratory for analyses of BTEX; TPH; alkalinity; moisture content; pH; iron; total phosphorous; total Kjeldahl nitrogen; and particle size analysis.

3.1.5 Soil Gas Permeability and Radius of Influence

A detailed description of the method for conducting a soil gas permeability test, including equations to compute k, the soil gas permeability, is given in the Test Plan and Technical Protocol (Hinchee et al., 1992).

Prior to air injection, the monitoring points were allowed to set up for 24 hours. A portable 1-HP explosion-proof positive displacement blower unit was used to inject air. After air injection was initiated, pressure readings were taken approximately every 1 to 2 minutes for the first hour, then approximately every 10 minutes for the following hour. The HyperventilateTM computer model was used to calculate the soil gas permeability.

3.2 Results and Discussion

3.2.1 Soil and Soil Gas Analyses

Results of the soil analyses for BTEX and TPH are presented in Table 7. The analytical report for this site is presented in Appendix B. All of the BTEX compounds were at concentrations below the detection limit in sample H1-VW-3'-3.5' and only small quantities of toluene (0.015 mg/kg) and total xylenes (0.051 mg/kg) were detected in sample H1-VW-4'-4.5'. TPH concentrations were low in sample H1-VW-3'-3.5' (12 mg/kg); however, 13,000 mg/kg of TPH was detected in sample H1-VW-4'-4.5'. The results of the soil chemistry analyses are summarized in Table 8.

Table 7. Results From Soil Analyses for BTEX and TPH at Building 1812

Sample Name	Benzene (mg/kg)	Toluene (mg/kg)	Ethylbenzene (mg/kg)	Total Xylenes (mg/kg)	TPH¹ (mg/kg)
H2-VW-3'-3.5'	< 0.062	< 0.072	< 0.052	< 0.093	12
H2-VW-4'-4.5'	< 0.0032	0.015	< 0.0026	0.051	13,000

Referenced to a reference oil composed of a mixture of 2,2,4-trimethylpentane, *n*-hexadecane, and chlorobenzene.

Table 8. Results From Soil Chemistry Analysis at Building 1812

	Sample Name				
Parameter	H2-	H2-VW-3'-3.5'		W-4'-4.5'	
Alkalinity (mg/kg CaCO ₃)		90		< 50	
Moisture (% by weight)		3.2		6.1	
pН	7.0			6.2	
Iron (mg/kg)	8,160			5,460	
Total Phosphorous (mg/kg)		450		600	
Total Kjeldahl Nitrogen (mg/kg)		66		53	
Particle Size Analysis (%)	Gravel:	10	Gravel:	1.0	
	Sand:	69.5	Sand:	76	
	Silt:	19	Silt:	21.5	
	Clay:	1.5	Clay:	1.5	

3.2.2 Soil Gas Permeability and Radius of Influence

The raw data for the soil gas permeability test at Building 1812 are presented in Appendix E. Using the HyperventilateTM computer model, soil gas permeabilities were calculated at each of the monitoring points. These data are presented in Table 9. The soil gas permeability varied considerably, with values ranging from 2.5 darcy up to 6.3 x 10° darcy. The radius of influence where 1 inch of water could be measured was calculated by plotting the log of the pressure change at the monitoring points versus the distance from the vent well (Figure 8). If pressure changes at all monitoring points are included, no radius of influence can be calculated based on these specifications. However, if the pressure change at monitoring points below the water table are not included (H2-MPA-5.0' and H2-MPB-5.0'), the radius of influence at Building 1812 is estimated to be approximately 7.5 feet.

3.2.3 Bioventing Demonstration

The decision was made to install a bioventing system at Building 1812. A 1-HP blower was installed on October 14, 1992. Air injection was initiated on October 14 at a flowrate of 4.25 scfm.

4.0 BACKGROUND AREA ACTIVITIES

The background area was located as shown in Figure 1. An existing monitoring well was used as the vent well and was screened from 1.5 feet to 9.0 feet. Soil samples were taken 2 feet from the monitoring well by hand auger. Site lithology at this area was representative of that in the contaminated areas.

A split-spoon soil sample was collected at a depth of 2.5 to 3.0 feet from the vent well borehole and was labeled H1-BKG-2.5'-3'. A soil vapor sample also was collected from the vent well after installation and labelled H1-BG-1.5-9. The soil samples were sent under chain of custody to Engineering-Science Inc., Berkeley Laboratory for analysis of BTEX; TPH; alkalinity; moisture content; pH; iron; total phosphorous; total Kjeldahl nitrogen; and particle size analysis. The soil vapor sample was sent under chain of custody to Air Toxics, Ltd., in Rancho Cordova, California, for analysis of BTEX and TPH.

Table 9. Results of Hyperventilate™ Soil Gas Permeability Analysis

Monitoring Point	Depth (ft)	Soil Gas Permeability (darcy)
H2-MPA	2.5	6.3 x 10°
	5.0	2.5
Н2-МРВ	2.5	3.1 x 10°
	5.0	430
	7.0	ND
H2-MPC	2.5	NR
	4.5	NR
	6.0	NR

ND No data were collected from this monitoring point.

NR No pressure readings were detected at this monitoring point.

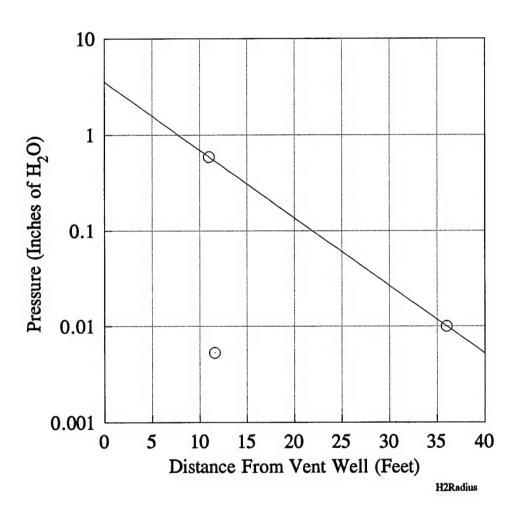


Figure 8. Radius of Influence at Building 1812

Results of the soil and soil vapor analyses for BTEX and TPH are presented in Table 10. The analytical report for the background area is presented in Appendix B. All of the BTEX compounds were at concentrations below the detection limit in the soil sample, and only a small quantity of TPH (84 mg/kg) was detected. The soil vapor sample contained higher concentrations of BTEX compounds with concentrations ranging from 0.16 ppmv (benzene) up to 4.7 ppmv (total xylenes). The results of the soil chemistry analyses also are summarized in Table 10.

An in situ respiration test was conducted at the background area beginning on October 9 after 24 hours of air injection. The test was concluded on October 11. Very little decrease in oxygen concentration occurred during the course of the in situ respiration test (Figure 9).

5.0 FUTURE WORK

Base personnel will be required to perform a simple weekly system check to ensure that the blower is operating within its intended flowrate, pressure, and temperature range. An on-site briefing for base personnel who will be responsible for blower system checks was conducted when the blowers were installed. The principle of operation was explained, and a simple checklist and logbook were provided for blower data. Base personnel will be asked to perform minor maintenance activities, such as replacing filters or gauges, or draining condensate from knockout chambers, but they will not be expected to perform complicated repairs or analyze gas samples. Replacement filters and gauges will be provided and shipped to the base, and serious problems, such as motor or blower failures, will be corrected by Battelle.

The progress of this system will be monitored by conducting semiannual respiration tests in the vent well and in each monitoring point and by regularly measuring the oxygen, carbon dioxide, and hydrocarbon concentrations in the extracted soil gas and comparing them to background levels. At least twice each year, the progress of the bioventing test will be reported to the base point-of-contact.

Table 10. Results From Soil Chemistry Analysis at Background Area

	Sample Name	
Parameter	Soil Sample H1-BKG-2.5'-3'	Soil Vapor Sample H1-BG-1.5-9
Benzene	<0.00060 mg/kg	0.16 ppmv
Toluene	<0.00070 mg/kg	0.93 ppmv
Ethylbenzene	<0.00050 mg/kg	0.43 ppmv
Total Xylenes	<0.00090 mg/kg	4.7 ppmv
ТРН	84 mg/kg	340 ppmv
Alkalinity (mg/kg CaCO ₃)	<50	
Moisture (% by weight)	9.8	
pН	6.6	
Iron (mg/kg)	11,600	
Total Phosphorous (mg/kg)	460	
Total Kjeldahl Nitrogen (mg/kg)	91	
Particle Size Analysis (%)	Gravel: 11.5	
	Sand: 48	
	Silt: 32	
	Clay: 8.5	

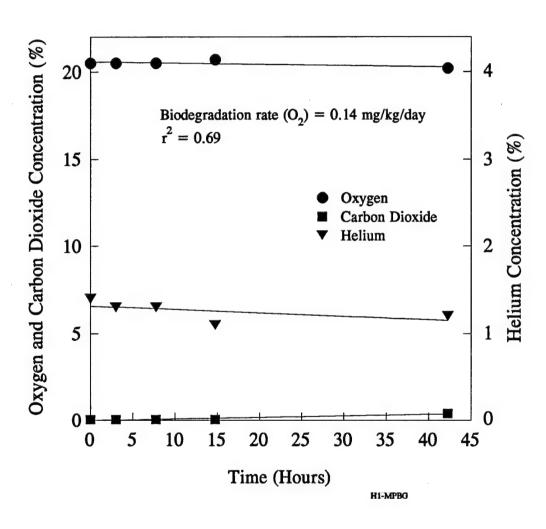


Figure 9. Oxygen Utilization and Carbon Dioxide Production During the In Situ Respiration Test at the Background Area

6.0 REFERENCE

Hinchee, R.E., S.K. Ong, R.N. Miller, D.C. Downey, and R. Frandt. 1992. *Test Plan and Technical Protocol for a Field Treatability Test for Bioventing* (Rev. 2), Report prepared by Battelle Columbus Operations, U.S. Air Force Center for Environmental Excellence, and Engineering Sciences, Inc. for the U.S. Air Force Center for Environmental Excellence, Brooks Air Force Base, Texas.

APPENDIX A

TEST PLAN FOR HANSCOM AFB, MASSACHUSETTS



505 King Avenue Columbus, Ohio 43201-2693 Telephone (614) 424-6424 Facsimile (614) 424-5263

September 2, 1992

Captain Catherine Vogel
Department of the Air Force
Building 1117
HQ AFESC/RDVW
Tyndall AFB, Florida 32403-6001

Dear Cathy:

SUBJECT: TEST PLAN FOR BIOVENTING INITIATIVE FIELD TEST AT BUILDING 1639 AND BUILDING 1812, HANSCOM AFB, MA.

This letter was prepared to accompany the report titled "Test Plan and Technical Protocol for a Field Treatability Test for Bioventing." The report was developed as a generic test plan for the Air Force Bioventing Initiative Project in which Hansom AFB is participating. This letter outlines site specific information to support the generic test plan.

The sites chosen for the bioventing test initiative are Building 1639 and Building 1812. Building 1639 is the base fuel service station (see map Figure 1). There have been three reported petroleum releases (gasoline, #2 fuel oil, and waste lubricating oil) at this facility since November 1990. The site at Building 1812 is the location of a heating oil fuel tank (see Figure 2). The tank was replaced recently and soil TPH concentrations as high as 2830 mg/Kg have been reported.

The purpose of this project is to investigate the feasibility of using the bioventing technology to remediate petroleum contaminated soils at the above mentioned sites.

Site Description:

Hanscom AFB is located in Bedford, Massachusetts. A generalized geologic and hydrogeologic column is shown in Figure 3. Groundwater at Hanscom is encountered at 3-8 feet. Figures 4 and 5 show test boring logs for two soil borings in the vicinity of the bioventing initiative sites.

Soil contamination at Building 1639 ranges from 599 ppm to 4400 ppm TPH. Soil samples at Building 1812 have indicated TPH concentrations as high as 2800 ppm. Boring and depth locations for these samples were not immediately available. Locations for initiating the soil gas survey at each site will be determined with direction from the Base POC.

Captain Catherine Vogel
Department of the Air Force
September 2, 1992
Page 2

Project Activities-

The following field activities are planned for the bioventing project at Hanscom AFB. The same procedures will be followed at each site. Additional detail can be found in Section 5.0 of the test plan and technical protocol.

- A small scale soil gas survey will be conducted to identify an appropriate location for installation of the bioventing system. The soil gas survey will be conducted in areas which site data have shown to be the most contaminated. Soil vapor from the candidate site should exhibit high petroleum hydrocarbon concentrations (10,000 ppm or greater), relatively low O₂ concentrations (0 % to 2.0 %), and relatively high CO₂ concentrations (depending on soil type, 2.0 % to 10.0 %, or higher). An uncontaminated background location will also be identified.
- Once the installation sites are located one vent well and three 3-level soil gas monitoring points will be installed in the contaminated location and one vent well will be installed in a background area (one background area will be used for both test sites, if possible). The wells and monitoring points will be installed using a two-man power auger or a portable drill rig to bore down to just above the water table. Three to four soil samples will be collected for chemical/physical analysis.
- 3- The air permeability test will be conducted in the contaminated test location.
- 4- Following the air permeability test, in situ respiration tests will be conducted in both the contaminated and the background test locations.
- Depending on the results of the air permeability test and the in situ respiration test, a decision will be made whether or not to install a blower system in the contaminated area for the long term bioventing test. If the decision is made to install the system, the blower will be plumbed to the vent well and bioventing will be started (assuming power is available). Site personnel will be trained for blower operation prior to Battelle leaving the site.
- A report detailing the results of the in situ respiration test and the air permeability test will be provided to the project officer and the base POC.

Schedule-

Field activities at Hanscom AFB are planned to begin on September 28, 1992. Battelle will have 2 to 3 people on site for approximately 3 weeks.

Captain Catherine Vogel Department of the Air Force September 2, 1992 Page 3

Base Support-

Hanscom AFB needs to be able to provide the following:

- Digging permits and utility clearance need to be obtained prior to the initiation of the field work. Underground utilities should be clearly marked to reduce the chance of utility damage or personal injury during soil gas probe and well installation. Battelle will not be able to begin field operations without these clearances.
- Electrical power will need to be easily accessible from the project site. The air permeability test and in situ respiration test can be performed using a gasoline powered electric generator. The operation of the bioventing system will require a permanent 220/110 V power source. If power will not be available immediately after the test is completed the bioventing system will be installed for start-up at a later date.
- The Air Force will need to provide drums to contain soil cuttings and provide for contaminated soil disposal.
- Base and site clearance will be required for Battelle's site employees. We will furnish you with personal information for each person at least one week prior to starting field operations.
 - Regulatory approval, if any is required, will need to be obtained by the base prior to start-up of the bioventing system. The system will likely be configured for air injection so there will be no point source vapor emission from the system. The wells to be installed will not intersect the apparent water table and no groundwater will be pumped. A letter from the Massachusetts DEP (Attachment 1) presented several concerns with the bioventing test plan. These concerns are addressed below.

Regulatory Issues-

- General Approval Requirements- DEP approval is necessary for the initiation of remediation of petroleum releases. This approval should be pursued by the base for the long-term bioventing test. The soil gas survey, vent well and monitoring point installation, air permeability test, and in situ respiration test are all standard field activities that generally do not require any regulatory approval prior to initiation. Battelle is available to discuss any of these activities with the DEP prior to start up of field activities.
- 2- Air Quality Permits and Approvals- The only air injection planned for this study is the injection of ambient atmospheric air into the vadose zone as a oxygen source for biological activity. Should conditions require the extraction of soil gas (i.e. a basement adjacent to the vent well) then treatment and permitting requirements will be discussed with the DEP.

Captain Catherine Vogel Department of the Air Force September 2, 1992 Page 4

- 3- Groundwater Remediation- The purpose of this research project is to investigate the efficacy of bioventing for the remediation of petroleum contaminated soils in the vadose zone. While bioventing, as configured for this project, does not directly address groundwater contamination, it has proven to be an effective technology for the remediation of petroleum contaminated soils. These soils typically act as the long-term source of localized groundwater contamination. Removal of the contaminant source is paramount in any groundwater remediation effort.
- 4- Nutrient Addition- Nutrient addition is not currently planned for this site.
- 5- Field Screening of Split-Spoon Samples- The Jar Headspace Technique will be employed for selected samples. Samples will be analyzed for BTEX via EPA Method 8020 and TPH via EPA Method 418.1. Samples will also be analyzed for nutrients.

Please let me know if there are any other regulatory concerns which need to be addressed. If you have any questions please feel free to call me at (614) 424-6122.

Sincerely,

Jeffrey A. Kittel
Researcher
Environmental Technology

JAK:mla

ATTACHMENT 1



Commonwealth of Massochusetts

Executive Office of Environmental Affairs

Department of Environmental Protection Metro Boston/Northeast Regional Office

William F. Weld Governor Daniel S. Greenbaum

JUN 1 0 1992

Mr. Robert Spelfogel Hanscom Air Force Base 3245th ABG/DEEV Bedford, MA 01730

RE: BEDFORD - Hanscom AFB
Bioventing Treatability Test
DEP Case #3-3882

Dear Mr. Spelfogel:

The Department of Environmental Protection is in receipt of a January 1992 "Report, Test Plan And Technical Protocol For A Field Treatability Test For Bioventing". This report was prepared for the U.S. Air Force by Battelle Corporation and was submitted to the Department by the Air Force with a request to provide information concerning the regulatory requirements of the bioventing remediation technique. Preliminary approval of the bioventing concept was also requested. Hanscom Air Force Base is planning to contract Battelle Corporation to conduct Bioventing Treatability Testing at Buildings 1812 and 1639.

Building 1639 (DEP Case #3-3882) is the Air Force Base Service Station. Department files indicate three petroleum release incidents at this location. These releases were assigned the incident response numbers N92-0837, N91-794 and N90-1884. Gasoline, #2 fuel oil, and waste lubricating oil have been released to soil and/or groundwater at this site.

At Building 1812 soil samples collected from borings were found to be contaminated with concentrations of Total Petroleum Hydrocarbons as high as 2830 mg/kg.

The bioventing process uses aeration of subsurface soils to stimulate in-situ biological activity and promote bioremediation. The following points outline the bioventing process.

Hanscom AFB Page 2

- (1) Forced air is supplied to the contaminated unsaturated zone to produce the aerobic conditions necessary for biodegradation.
- (2) Airflow is optimized to reduce volatilization while maintaining aerobic conditions.
- (3) Local soil gas conditions are monitored to assure aerobic conditions.
- (4) Moisture and nutrients are added to the contaminated unsaturated zone if necessary.
- (5) Dewatering is conducted, if necessary to lower the water table below the contaminated soils.

After review of the above mentioned report the Department hereby approves of the initiation of Bioventing Field Treatability Testing at Buildings 1639 and 1812. The following comments are also provided:

* General Approval Requirements.

Departmental approval must be obtained for the initiation of the remediation of releases of petroleum or becardous materials as locations To Be Investigated or

hazardous materials at Locations To Be Investigated or Confirmed Disposal Sites. A bioventing system to remediate petroleum or hazardous materials releases at Hanscom Air Force Base is subject to the above-mentioned approvals.

* Air Quality Permits and Approvals.

Page 5 Section 2.1.3 states that when air is injected into a contaminated zone and withdrawn from clean soils volatile hydrocarbons are allowed to degrade prior to being withdrawn thereby eliminating contaminated off-gases. "This configuration typically does not require air emission permitting".

Be advised that all air emissions resulting from treatment systems at petroleum or hazardous materials disposal sites are subject to the Air Pollution Control Regulations at 310 CMR 6.00-8.00 in addition to the requirements of the Massachusetts Contingency Plan at 310 CMR 40.000.

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Hanscom AFB Page 3

> In general, if an air contaminant source emits less than one ton of contaminants per year, then that emission source is not subject to application for approval from the Department of Environmental Protection's Division of Air Quality.

Whether or not the emission source emits more than one ton of pollutants per year, if the emission is part of a system being operated to remediate a release of petroleum or hazardous materials to the environment, then that system is subject to M.G.L. c.21E and the MCP. The DEP Bureau Of Waste Site Cleanup requires that off-gas controls be installed on all point source remedial air emissions where Soil Vapor Extraction systems are utilized. Off-gas controls must be implemented during the first 180 days of operation at a minimum. Off-gas treatment must reduce vapor-phase emissions of VOC's by at least 95 percent. Approval to operate without emission controls after the first 180 days are contingent upon receipt of information confirming the absence of a threat of harm to human health or the environment.

Page 5 indicates that during a 1988 study at Hill Air Force Base it became apparent that bioventing had great potential for remediating JP-4 fuel-contaminated soils.

It is not clear however that bioventing is effective for remediation of groundwater contaminated with fuel oils. Many of the components contained in aircraft fuels, diesel fuel and heavier petroleum fuel products are not significantly volatile and would not readily be transferred from the groundwater to the unsaturated zone where they could be treated by the bioventing system. In order to achieve a permanent solution to the petroleum contamination present at a site it may be necessary to reduce the levels of contaminants in soil and groundwater to levels which do not pose a significant or otherwise unacceptable risk to public health, safety, public welfare or the environment.

Page 6 indicates that adding nutrients as required to increase biodegradation rates may be necessary. The addition of nutrients to the subsurface is subject to the requirements of M.G.L. c.21 and the Groundwater Discharge Permit Regulations (314 CMR 5.00). Nutrient addition may be initiated only upon receipt of a Groundwater Discharge Permit issued by the DEP Division of Groundwater Pollution Control.

Hanscom AFB Page 4

Page 42, Section 2.1.2 - Exploratory Boring in Deep Soils, states that "Split-spoon samples will be visually checked for fuel contamination and screened for volatile emissions by passing a hydrocarbon analyzer slowly over the open split-spoon".

This method is not acceptable for the field screening of split spoon samples for petroleum hydrocarbon contamination. The Jar Headspace Technique (Reference: "Management Procedures for Excavated Soils Contaminated with Virgin Petroleum Oils", Policy #WSC-89-001) should be applied for the measurement of volatile petroleum constituents. Total Petroleum Hydrocarbons by Infrared (Standard Methods 503 or 5520) or Oil Fingerprinting (ASTM D 3328) should be used to determine concentrations of non-volatile petroleum hydrocarbons.

If you have any questions regarding this matter please do not hesitate to contact Jack Miano at the letterhead address or telephone (617) 935-2160 X142.

Very truly yours,

Jack Miano

Environmental Engineer

Stephen M. Johnson

Acting Chief,

Site Management Branch

SMJ/JM/ae

CC: DEP, BWSC, Boston, Attn: Jeff Krukonis DEP, DWS, NERO, Attn: Jim Persky

Bedford BOH, Attn: David Black

Bedford DPW, 312 Great Rd., Bedford, MA 01730,

Attn: Mr. Richard Warrington, Director

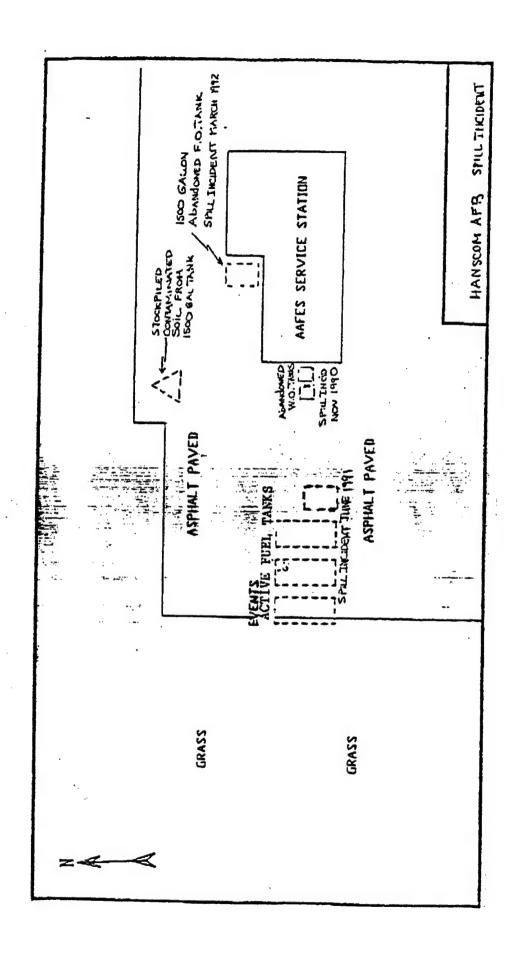


FIGURE 1. SITE DIAGRAM FOR BUILDING 1639

FORMER LOCATION OF UNDERGROUND STORAGE TANK BUILDING NO. 1816

HANSCOM AIR FORCE BASE, MASSACHUSETTS

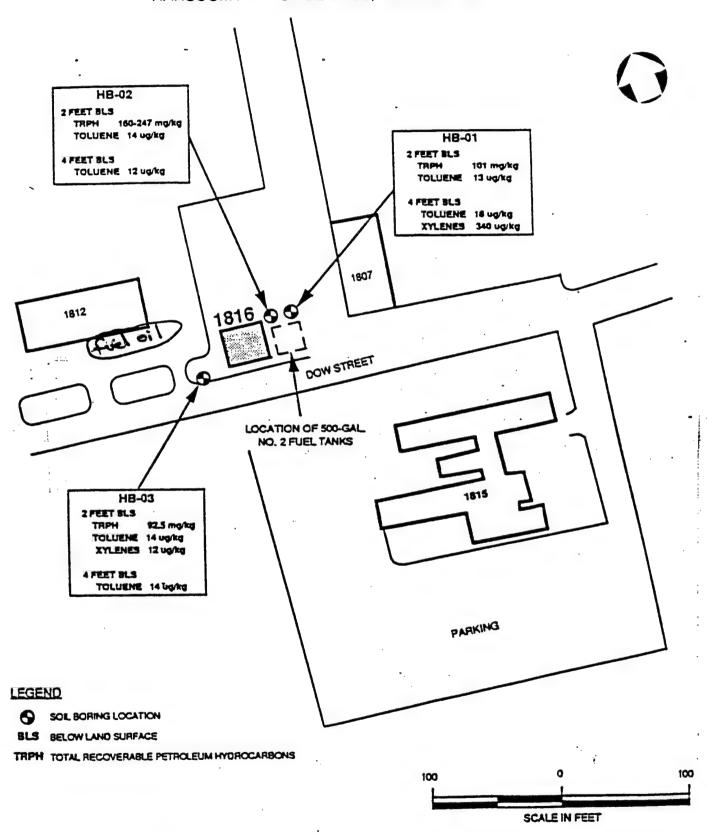


FIGURE 2. SITE DIAGRAM FOR BUILDING 1812

GENERALIZED GEOLOGIC AND HYDROGEOLOGIC COLUMN INVESTIGATION OF SUSPECTED HAZARDOUS WASTE SITES HANSCOM AIR FORCE BASE, MASSACHUSETTS

PERIOD	ЕРОСН	FORMATION	COLUMNAR SECTION	APPROXIMATE THICKNESS	CHARACTER	
	RECENT	FILL —unconformable—		0-10	TOPSOIL & CLEAN SANDY FILL	
		PEAT —unconformable—		0-7	BLACK ORGANIC SANDS & PEAT	
-	(NIS	? OUTWASH DEPOSITS	منہ	0-18	Clean Coarse to fine Sands SATURATED, PERMEABLE	UPPER AQUIFER
QUATERNARY	PLEISTOCENE (WISCONSIN)	LACUSTRINE DEPOSITS		0-65	Fine SANDS & SILTS Grading to dayey Silts, Low permeability, SATURATED	LACUSTRINE AQUITARD
		Sub-Lacustrine Deposits —unconformable—	مم	0-20	Fine Sands & Silts, Saturated Permeable	
		GLACIAL TILL —unconformable—		0-10	Coarse to fine SAND, trace to some SILT, trace to some GRAVEL	LOWER AQUIFER
		ANDOVER GRANITE		0-?	Medium to Coarse Grained Muscovite- Biotite Granite	BEDROCK AQUIFER

BORING NUMBER	HB-19	4
JOB NUMBER	11-9540	
DATE STARTED	4-24-90	
DATE COMPLETED	4-24-90	
DRILLED BY	TN	
LOGGED BY	KMP	
CHECKED BY	GPM	

REMARKS:

PAGE 1 OF 1

FLUSH MOUNTED WELL SWL MEASURED 9.34' BELOW TOC ON 5/29/90 WELL ALIGNMENT TEST COMPLETED 5/18/90 HNU READINGS RECORDED IN "LAB TEST" COLUMN

ELEV. IN FEET	DEPTH IN FEET	DESCRIPTION	MONITORING WELL CONSTRUCTION	SYM- BOLS	LAB TESTS	SPT N VALUE
129.02	0.0	Firm yellowish brown (10 YR 5/6) to very pale brown (10 YR 7/3) fine to medium grained SAND (SP)			0	19
122.02	7.0				0	22 -
		Firm light brownish gray (2.5 YR 6/2) fine grained SAND (SP)			. 0	17
120.02		Very stiff light yellowish brown (2.5 Y 6/4) SILT (ML)		<u></u>	0	16 -
117.02	12.0	Very stiff light yellowish brown (2.5 Y 6/4) SILT with lenses of fine grained sand (ML)				· 17
113.52	15.5	Boring Terminated at 15.5				16 -
-						
		<i>,</i>				
		·				
+						-
1						

BORING NUMBER	HB-03
JOB NUMBER	11-9540
DATE STARTED	5-1-90
DATE COMPLETED	5-1-90
ORILLED BY	TN
LOGGED BY	KMP
CHECKED BY	GPM

REMARKS:	
HEIMAHAS.	

PAGE 1 OF 1

FLUSH MOUNTED WELL SWL MEASURED 3.05 BELOW TOC ON 5/29/90 WELL ALIGNMENT TEST COMPLETED 5/18/90 HNU READINGS RECORDED IN "LAB TEST" COLUMN

F	LEV. IN EET	DEPTH IN FEET	DESCRIPTION	MONITORING WELL CONSTRUCTION	SYM- BOLS	LAB TESTS	SPT N VALUE
13	32.00	0.0	Very stiff to hard light olive brown (2.5 Y 5/4) SILT (ML)			•	
		·			<u>*</u>	0	32
						0	20 —
12	3.00	9.0	Firm vallewish has we (40 VC 540 5			0	27
12	0.00	12.0	Firm yellowish brown (10 YR 5/6) fine to medium grained SAND with stratifications (SM)			0	17 -
		13.0				0	31
1		,	Boring Terminated at 12.0°				
							
							4
:	.						-
_							

APPENDIX B

ANALYTICAL REPORT FOR BUILDING 1639, BUILDING 1812, AND BACKGROUND AREA

(a) AIR TOXICS LTD.

AN ENVIRONMENTAL ANALYTICAL LABORATORY

WORK ORDER #: 9210075

Work Order Summary

CLIENT:

Mr. Jeff Kittel

BILL TO:

Accounts Payable

Battelle

Engineering Science

505 King Ave.

1700 Broadway Ste. 900

Columbus, OH 43201

Denver, CO 80290

PHONE:

614-424-6122

INVOICE # 8630

FAX:

614-424-3667

P.O. # DE268.03.04

DATE RECEIVED:

AMOUNT: \$910.00

DATE REPORTED:

10/14/92 10/26/92

PROJECT # G4468-0640

FRACTION #	NAME	TEST	VAC./Press.	PRICE
01A	H1-A-2.5	TO-3	0 "Hg	\$120.00
02A	H1-A-5	TO-3	0 "Hg	\$120.00
03A	H1-B-2.5	TO-3	2.0 "Hg	\$120.00
04A	H1-B-5	TO-3	1.0 "Hg	\$120.00
05A	H1-C-3.5	TO-3	0.5 "Hg	\$120.00
06A	H1-C-6	TO-3	0.5 "Hg	\$120.00
07A	H1-BG-1.5-9	TO-3	1.0 "Hg	\$120.00
07B	H1-BG-1.5-9 Duplicate	TO-3	1.0 "Hg	NC
08A	Method Spike	TO-3	NA	NC
09A	Lab Blank	TO-3	NA	NC
Misc Charges	1 Liter SUMMA Canister P.	reparation (7)	@ \$10.00 each.	\$70.00

CERTIFIED BY Mola & Frances

Laboratory Director

SAMPLE NAME: H1-A-2.5 ID#: 9210075-01A

EPA Method TO-3

(Aromatic Volatile Organics in Air)

BTXE BY GC/PID

File Name: Dil, Factor:	610150		Date of Collection	
Dit. Pactor.	MDL	MDL	Date of Analysis: Amount	10/15/92 Amount
Compound	(ppmv)	(uG/L)	(ppmv)	(uG/L)
Benzene	0.20	0.62	5.2	16
Toluene	0.20	0.74	4.2	15
Total Xylenes	0.20	0.85	2.9	12
Ethyl Benzene	0.20	0.85	1.1	4.7

TOTAL PETROLEUM HYDROCARBONS GC/FID

File Name:	610150	8	Date of Collection	1: 10/12/92
Dil. Factor:	20	0	Date of Analysis:	10/15/92
_	MDL	MDL	Amount	Amount
Compound	(ppmv)	(uG/L)	(ppmv)	(uG/L)
TPH*	2.0	8.0	5600	22000

^{*}TPH referenced to Jet Fuel (MW=156)

SAMPLE NAME: H1-A-5 ID#: 9210075-02A

EPA Method TO-3

(Aromatic Volatile Organics in Air)

BTXE BY GC/PID

File Name: Dil. Factor:	610150 50		Date of Collection Date of Analysis:	
	MDL	MDL	Amount	Amount
Compound	(ppmv)	(uG/L)	(ppmv)	(uG/L)
Benzene	0.50	1.6	27	84
Toluene	0.50	1.8	35	130
Total Xylenes	0.50	2.1	30	130
Ethyl Benzene	0.50	2.1	10	42

TOTAL PETROLEUM HYDROCARBONS GC/FID

File Name: 6101509 Date of Collection: 10/12/92 Dil. Factor: 500 Date of Analysis: 10/15/92					
Compound	MDL (22222)	MDL (C (L)	Amount	Amount	
Compound	(ppmv)	(uG/L)	(ppmv)	(uG/L)	
TPH*	5.0	20	19000	76000	

^{*}TPH referenced to Jet Fuel (MW=156)

SAMPLE NAME: H1-B-2.5 ID#: 9210075-03A

EPA Method TO-3

(Aromatic Volatile Organics in Air)

BTXE BY GC/PID

File Name:	610151	0	Date of Collection	ı: 10/12/92
Dil. Factor:	22	0	Date of Analysis:	10/15/92
	MDL	MDL	Amount	Amount
Compound	(ppmv)	(uG/L)	(ppmv)	(uG/L)
Benzene	0.22	0.69	2.8	8.7
Toluene	0.22	0.81	1.3	4.8
Total Xylenes	0.22	0.93	1.9	8.1
Ethyl Benzene	0.22	0.93	0.84	3.6

TOTAL PETROLEUM HYDROCARBONS GC/FID

File Name:	610151	0	Date of Collection	ı: 10/12/92
Dil. Factor:	22	O	Date of Analysis:	10/15/92
	MDL	MDL	Amount	Amount
Compound	(ppmv)	(uG/L)	(ppmv)	(uG/L)
TPH*	2.2	8.8	2700	11000

^{*}TPH referenced to Jet Fuel (MW=156)

SAMPLE NAME: H1-B-5 ID#: 9210075-04A

EPA Method TO-3

(Aromatic Volatile Organics in Air)

BTXE BY GC/PID

File Name:	610160	16	Date of Collection	n: 10/12/92
Dil. Factor:	21	0	Date of Analysis:	10/16/92
	MDL	MDL	Amount	Amount
Compound	(ppmv)	(uG/L)	(ppmv)	(uG/L)
Benzene	0.21	0.66	2.4	7.5
Toluene	0.21	0.77	0.84	3.1
Total Xylenes	0.21	0.89	1.5	6.4
Ethyl Benzene	0.21	0.89	0.42	1.8

TOTAL PETROLEUM HYDROCARBONS GC/FID

File Name;	610160	16	Date of Collection	ı: 10/12/92
Dil. Factor:	21	0	Date of Analysis:	10/16/92
	MDL	\mathbf{MDL}	Amount	Amount
Compound	(ppmv)	(uG/L)	(ppmv)	(uG/L)
TPH*	2.1	8.4	3200	13000

^{*}TPH referenced to Jet Fuel (MW=156)

SAMPLE NAME: H1-C-3.5 ID#: 9210075-05A

EPA Method TO-3

(Aromatic Volatile Organics in Air)

BTXE BY GC/PID

File Name:	610160	7	Date of Collection	
Dil. Factor:	1	0	Date of Analysis:	10/16/92
	MDL	MDL	Amount	Amount
Compound	(ppmv)	(uG/L)	(ppmv)	(uG/L)
Benzene	0.010	0.031	0.44	1.4
Toluene	0.010	0.037	0.13	0.48
Total Xylenes	0.010	0.042	0.37	1.6
Ethyl Benzene	0.010	0.042	0.11	0.47

TOTAL PETROLEUM HYDROCARBONS GC/FID

File Name: Dil. Factor:	610160	17 0	Date of Collection Date of Analysis:	
Compound	MDL (ppmv)	MDL (uG/L)	Amount (ppmv)	Amount (uG/L)
TPH*	0.10	0.40	280	1100

^{*}TPH referenced to Jet Fuel (MW=156)

SAMPLE NAME: H1-C-6 ID#: 9210075-06A

EPA Method TO-3

(Aromatic Volatile Organics in Air)

BTXE BY GC/PID

File Name:	610151		Date of Collection	
Dil. Factor:	MDL MDL	WDL	Date of Analysis: Amount	10/15/92 Amount
Compound	(ppmv)	(uG/L)	(ppmv)	(uG/L)
Benzene	2.1	6.6	11	34
Toluene	2.1	7.7	20	74
Total Xylenes	2.1	8.9	67	2 80
Ethyl Benzene	2.1	8.9	9.3	39

TOTAL PETROLEUM HYDROCARBONS GC/FID

File Name:	610151	4	Date of Collection	a: 10/12/92
Dil. Factor:	210	0	Date of Analysis:	10/15/92
	MDL	MDL	Amount	Amount
Compound	(ppmv)	(uG/L)	(ppmv)	(uG/L)
TPH*	21	84	11000	44000

^{*}TPH referenced to Jet Fuel (MW=156)

SAMPLE NAME: H1-BG-1.5-9 ID#: 9210075-07A

EPA Method TO-3

(Aromatic Volatile Organics in Air)

BTXE BY GC/PID

File Name: Dil. Factor:	610160 1)4 0		te of Collection: 10/12/92 te of Analysis: 10/16/92	
	MDL	MDL	Amount	Amount	
Compound	(ppmv)	(uG/L)	(ppmv)	(uG/L)	
Benzene	0.010	0.031	0.16	0.50	
Toluene	0.010	0.037	0.93	3.4	
Total Xylenes	0.010	0.042	4.7	20	
Ethyl Benzene	0.010	0.042	0.43	1.8	

TOTAL PETROLEUM HYDROCARBONS GC/FID

File Name:	610160	4	Date of Collection	ı: 10/12/92
Dil. Factor:		0	Date of Analysis:	10/16/92
	MDL	MDL	Amount	Amount
Compound	(ppmv)	(uG/L)	(ppmv)	(uG/L)
TPH*	0.10	0.40	330	1300

^{*}TPH referenced to Jet Fuel (MW=156)

SAMPLE NAME: H1-BG-1.5-9 Duplicate ID#: 9210075-07B

EPA Method TO-3

(Aromatic Volatile Organics in Air)

BTXE BY GC/PID

File Name:	610160	5	Date of Collection	1: 10/12/92
Dil, Factor:	1	0	Date of Analysis:	10/16/92
	MDL	MDL	Amount	Amount
Compound	(ppmv)	(uG/L)	(ppmv)	(uG/L)
Benzene	0.010	0.031	0.16	0.50
Toluene	0.010	0.037	0.93	3.4
Total Xylenes	0.010	0.042	4.7	20
Ethyl Benzene	0.010	0.042	0.43	1.8

TOTAL PETROLEUM HYDROCARBONS GC/FID

File Name: Dil. Factor:	610160 1		Date of Collection Date of Analysis:	•
	MDL	MDL	Amount	Amount
Compound	(ppmv)	(uG/L)	(ppmv)	(uG/L)
TPH*	0.10	0.40	340	1400

^{*}TPH referenced to Jet Fuel (MW=156)

SAMPLE NAME: Method Spike ID#: 9210075-08A

EPA Method TO-3

(Aromatic Volatile Organics in Air)

BTXE BY GC/PID

File Name: Dil. Factor:	610160 1)2 0	Date of Collection: 10/12/92 Date of Analysis: 10/16/92
	MDL	MDL	Date of Analysis: 10/10/92
Compound	(ppmv)	(uG/L)	% Recovery
Benzene	0.001	0.003	90
Toluene	0.001	0.004	83
Total Xylenes	0.001	0.004	80
Ethyl Benzene	0.001	0.004	80

TOTAL PETROLEUM HYDROCARBONS GC/FID

File Name: Dil. Factor:	610160	3 0	Date of Collection: 10/12/92
	MDL	MDL	Date of Analysis: 10/16/92
Compound	(ppmv)	(uG/L)	% Recovery
TPH*	0.010	0.040	110

^{*}TPH referenced to Jet Fuel (MW=156)

SAMPLE NAME: Lab Blank ID#: 9210075-09A

EPA Method TO-3

(Aromatic Volatile Organics in Air)

BTXE BY GC/PID

File Name: Dil. Factor:	610150 1.		Date of Collection	**************************************
DH. FRCUIT.	MDL	MDL	Date of Analysis: Amount	Amount
Compound	(ppmv)	(uG/L)	(ppmv)	(uG/L)
Benzene	0.001	0.003	Not Detected	Not Detected
Toluene	0.001	0.004	Not Detected	Not Detected
Total Xylenes	0.001	0.004	Not Detected	Not Detected
Ethyl Benzene	0.001	0.004	Not Detected	Not Detected

TOTAL PETROLEUM HYDROCARBONS GC/FID

File Name:	610150	1	Date of Collection	n: 10/12/92
Dil. Factor:	1.	0	Date of Analysis:	10/15/92
	MDL	MDL	Amount	Amount
Compound	(ppmv)	(uG/L)	(ppmv)	(uG/L)
TPH*	0.010	0.040	Not Detected	Not Detected

^{*}TPH referenced to Jet Fuel (MW=156)



CHAIN OF CUSTODY RECORD

SUITE	CA 95743	638-9917
SOUNDS GOLD CIRCLE, SUITE 'F	RANCHO CORDOVA, CA 9574	(916) 638-9892 • FAX (916) 638-9917
SE GOLD	SCHO CC	3-9892 • F
SOUNTE	HA	(916) 638

	OF COSTODI NECOKD	Page of	_
PROJECT # (5 4468 - 0640_PO#	COLLECTED BY (Signature) MAT / TA		
FIELD SAMPLE 1.D.# SAMPLING MEDIA (Tenax, Canister etc.) 41-A-2.5 41-A-5 11 41-B-2.5 11 41-B-3.5 11 41-C-6 11 41-89-7.59 11	DATE/TIME ANALYSIS 12.0.cf / 09.00 B.TEX / TV.H 11	VAC./PRESSURE LAB I.D. # 0."/# 2.0"/# 2.0"/# 0.5"/#	#

RECEIVED BY: DATE/TIME	CONDITION	
RELINQUISHED BY: DATE/TIME	TEMP(°C)	
IVED BY: DATE/TIME RELIN	LAB USE ONLY OPENED BY: DATE/TIME	
RECE	AIR BILL #	
RELINQUISHED BY: DATE/TIME	SHIPPER NAME	REMARKS

BERKELEY LABORATORY 600 BANCROFT WAY BERKELEY, CA 94710 Tel: (415) 841-7353

Report Date: November 19, 1992

Work Order No.:4432

Client:

Doug Downey

ES Denver/AFCEE/Hanscom AFB

1700 Broadway Denver, CO 80290

Date of Sample Receipt: 10/05/92

Your soil samples identified as:

H1-VW-4'-4.5' H1-VW-4.5'-5.0' H1-A-3'-4' H2-VW-3'-3.5' H2-VW-4'-4.5' H1-BKG-2.5'-3'

were analyzed for BTEX by EPA Method 8020, TRPH by EPA Method 418.1, TKN, total phosphorus, soil classification, pH, alkalinity, iron and moisture.

The analytical reports for the samples listed above are attached.

GC VOLATILES DATA PACKAGE

VOLATILE ORGANICS CASE NARRATIVE WORK ORDER NO. 4432 EPA METHOD 8020

Sample H1-VW-S³-3.5' (4432-04) was run as a medium level due to the presence of high concentrations of non-target analytes.

GC AMALYTICAL REPORT Analytical Dethou BTEX Aromatic Compounds

Work Order No.:4452

Client ID:HI-UW-47-4.57

% Moisture: 22.21

Matrix: SUIL

Laboratory ID:4452-1

Level:LOW

Date Collected: 10/02/92

Unit:UG/KG

Dilution Factor:

Date Analyzed:10/15/92 Date Confirmed:NA

Compound	Result	Reporting Limit	
			.: :::
Benzene	48.0 D=5	0.0	
Ethyl Benzene	17.0	Ü.o	
for logging	20.0	0.9	
Xylenes (total)	20.0	1.2	

ND-Not Detected NA-Not Applicable D-Dilution Factor

ANALYST: AS

GROUP LEADER: hur

GC ANALYTICAL REPORT Analytical Method BIEX Aromatic Compounds

Work Order NO.:4432

% Moisture: 21.81

Client ID:HI-UW-4,5/-5.0/

Matrix: SUIL

Laboratory ID:4432-2

Level:MEDIUM

Date Collected: 10/02/92

Unit:UG/KG

Dilution Factor:

Date Analyzed:10/14/92 Date Confirmed:NA

Compound	Result	Reporting Limit
Benzene	670.Ú	77.0
Ethyl Benzene	450.0	64.0.
Tolhene	270.0	90.0
Xvlenes (total)	450.0	120.0

ND-Not Detected NA-Not Applicable D-Dilution Factor

ANHLYST: AB

GROUP LEADER : Lugar

GC ANALYTICAL REPORT Analytical Method BTEX Aromatic Compounds

Work Order NO.:4432

Client ID:HI-A-3'-4'

Laboratory ID:4432-3

Date Collected: 10/02/92

Dilution Factor:

_

% Moisture: 5.81

Matrix:SOIL

Level:MEDIUM

Unit:UG/KG

Date Analyzed:10/14/92

Date Confirmed:NA

Campound	Result	Reporting Limit	
Benzene	1000.0	130.0	
Ethyl Benzene	1300.0	110.0	
Toluene	4500.0	150.0	
Xylenes (total)	12000.0	190.0	

ND-Not Detected NA-Not Applicable D-Dilution Factor

ANALYST: #3

GROUP LEADER: hour

GC ANALYTICAL REPORT Hnalutical Method BTEX Aromatic Compounds

Order NO.:4432 Client ID:HZ-VW-3:-3.5: Work Order NO.:4432

% Moisture: 3.2

Matrix: SUIL

Laboratory 1D:4432-4

Level:MEDIUM

Date Collected: 10/03/92

Unit:UG/KG

Dilution Factor:

Date Analyzed:10/14/92 Date Confirmed:NA

Campound	Result	Reporting Limit	
Benzene	NÚ	52.0	
Ethyl Benzene	· ND	52.0	
Toluene	ND	72.0	
Xylenes (total)	ND	93.0	

ND-Not Detected NA-Not Applicable D-Dilution Factor

ANHLYST: AS

GROUP LEADER: Kuse

GC ANALYTICAL REPORT Analytical Method BTEX Aromatic Compounds

Work Order No. : 4452 TP 11/18/92

Client ID: H2-UW-4/-4.5/

% Moisture: 6.14

Matrix:50IL

Level:LOW

Unit:UG/KG

Date Collected: 10/05/92

Laboratory ID:4452-5

Dilution Factor:

Į-

Date Analyzed:10/15/92

Date Confirmed:NA

Compound	Result	Reporting Limit
Kenzene	ND	3.2 ·
Ethyl Benzene	ИD	Z.6
ividene	15.0	3.7
Xylenes (total)	51.0	4.8

ND-Not Detected NA-Not Applicable D-Dilution Factor

ANALYST # AB

GROUP LEADER: Rever

GC ANALYTICAL REPORT Hnalutical Method BIEX Aromatic Compounds

Work Order NO.:4432

Client ID:HI-BkG-2.5/-5/

% Moisture:NA

Matrix: SOIL

Laboratory ID:4432-7

Level:LOW

Date Collected: 10/04/92

Unit:UG/KG

Dilution Factor: 1

Date Analyzed:10/13/92

Date Confirmed:NA

Compound	Result	Reporting Limit
=======================================		
Benzene	ИD	0.6
Ethyl Benzene	Nū	0.5
loluene	ND	0.7
Xylenes (total)	ND	0.9

ND-Not Detected NA-Not Applicable D-Dilution Factor

ANALYST: AD

GROUP LEADER: MON

GC ANALYTICAL REPORT Analytical Method BIEX Aromatic Compounds

Work Order NO.:4432

Client ID:METHOD BLANK

Laboratory ID:MSUG4921015

Date Collected: NA

Dilution Factor:

% Moisture:NA

Matrix:SOIL

Level:LOW

Unit:UG/KG

Date Analyzed:10/15/92

Date Confirmed:NA

Compound	Result	Reporting Limit	
			M 12 10 101
Benzene	NO	0.6	
Ethul Benzene	ND	0.5	
Toluene	พีบี.	0.7	
Xylenes (total)	ND	0.9	,

ND-Not Detected NA-Not Applicable D-Dilution Factor

ANALYST: AD

GROUP LEADER: fund

GC ANALYTICAL REPORT Hnalutical Method BIEX Aromatic Compounds

Work Order NO.:4452

% Moisture:NA

Client ID:METHOD BLANK

Matrix: SOIL

Laboratory ID:MWVG4921014

Level:MEDIUM

Date Collected: NA

Unit:UG/KG

Dilution Factor: 1

Date Analyzed:10/14/92

Date Confirmed:NA

Compound	Result	Reporting Limit
Benzene	НŪ	60.0
Ethyl Benzene	ND ·	90.0
toluena	ND	70.0
Xvlenes (total)	ND	90.0

ND-Not Detected NA-Not Applicable D-Dilution Factor

ANALYST: AS

GC ANALYTICAL REPORT Analytical Method BTEX Aromatic Compounds

Work Order No.:4452

Client ID: METHOD BLANK

Laboratory ID:M5V64921015

Date Collected: NA

Dilution Factor: 1 % Moisture:NA

Matrix: SUIL

Level:LOW

Unit:UG/KG

Date Analyzed:10/15/92 Date Confirmed:NA

(Jámpound	Result	Reporting Limit	
F.	Benzene	ND		0.6
	Ethyl Benzene	ND		0.5
7	Toluene	ND		0.7
>	(ylenes (total)	ND		0.9

ND-Not Detected NA-Not Applicable D-Dilution Factor

ANALYST : 16

GROUP LEADER: hun

FS-ENGINEERING SCIENCE, INC.

LABORATORY NO.

600 BANCROFT WAY BERKELEY, CA 94710

GC ANALYTICAL REPORT ANALYTICAL REPORT BTEX AROMATIC COMPOUNDS

MATRIX: SOIL

DATE:10/14/92

CLIENT ID a-a-a-TriFluoro

Tuluene

MWUG4921014	METHOD BLANK	. yl
4452-2	HI-UW-4.57-5.07	ó 2
4432-3	H1-A-3'-4'	115
4452-4	HVW-5'-3.5'	1 Z Z
	+p 11/1x	

LABORATORY NO.

600 BANCROFT WAY BERKELEY, CA 94710

GC ANALYTICAL REPORT -ANALYTICAL REPORT BTEX AROMATIC COMPOUNDS

MATRIX: SOIL

DATE:10/13&15/92

a-a-a-TriFluoro

	Toluene				
MSUG4921013	METHOD BLANK	94			
4432-7	HI-BKG-2.5'-3'	114			
4435-1	KAFB66-SB1-SS3-5-5.51	108			
MSVG4921015	METHOD BLANK	79			
4432-1	HI-UW-4'-4.5'	118			
4452-1DIL	HI-VW-41-4.51DIL	104			
4432-5	HF-UW-41-4.51	20			
4451-1	H	109			
	TP 11/18				

CLIENT ID

QUALITY CONTROL RESULTS SUMMARY ANALYTICAL REPORT BTEX AROMATIC COMPOUNDS

Work Order No.: 4432,4451,4455

QC sample No.: 55VG4921009A&B

Date analyzed:10/09/92

Matrix: SOIL

Dilution factor:1

:			MSD MSD PR UG/KG	RPD	QC LIMITS RPD PR
COMPOUND 8020 analysis	 S# SR UG/KG UG/KG	MS MS FR UG/KG	MSD MSD PR UG/KG	RPD	QC LIMITS RPD PR
Benzene Toluene 	20 ND 20 ND 1	1	23.5 118 	İ	29 39-150

MS = Spike sample

MSD = Spike sample duplicate

SH = Spike added

SR = Sample result

ND = Not Found At or Above Detection Limits

 $RPD = 100 \times (MS-MSD)/((MS+MSD)/2)$

 $PR = 100 \times ((MS \text{ or } MSD) - SR)/SA$

ANALYST: AT

ac: My

= Not calculated

= Not Applicable

= Out of limits

NC

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QUALITY CONTROL RESULTS SUMMARY ANALYTICAL REPORT BTEX AROMATIC COMPOUNDS

Work Order No.: 4432,4451,4435

QC sample No.: SWVG4921016A&B Date analyzed:10/16/92

Matrix: SOIL Dilution factor:1

*========											
	onno Uysis	 SA UG∕KG	 SR UG/KG	•	MS PR 	 MSD UG/KG	•	 RPD 	IQC LI I IRPD	IMITS	*
COMF 8020 ana	OUND lysis	 SA UG/KG	 SR UG/KG	•	IMS IPR	I I MSD IUG/KG	•	I IRPD	IQC LI I IRPD	IMITS PR	*
Benzene Toluene 		İ	Ì	2000 2600 	İ	1640 2700	82 135 	1	İ	39-150 - 46-148 	*

MS = Spike sample

MSD = Spike sample duplicate

SR = Sample result

SA = Spike added

ND = Not Found At or Above Detection Limits

 $RPD = 100 \times (MS-MSD)/((MS+MSD)/2)$

 $PR = 100 \times ((MS \text{ or } MSD) - SR)/SA$

ANALYST: A

ac: pws

= Not calculated

= Not Applicable

= Out of limits

NC

NA

WO # 4432,4451,4435

LAB NAME : ENGINEERING-SCIENCE, INC. DATE ANALYZED 10/13&15/92

LAB SAMPLE ID:MSUG4921013&15

DATE EXTRACTED : NA

MATRIX :LOW SOIL

INSTRUMENT ID: VGC-4

LAB	CLIENT	DATE
SAMPLE ID	SAMPLE ID	ANALYZED
MSVG4921013 4432-7 4435-1 MSVG492101 4432-1 4432-1DIL 4432-5 4451-1	METHOD BLANK HI-BKG-2.5'-3' KAFB66-SB1-SS3-5-5.5' METHOD BLANK HI-UW-4'-4.5' HI-UW-4'-4.5' DIL HF-UW-4'-4.5' MC-UW-7	10/13/92 10/13/92 10/13/92 10/15/92 10/15/92 10/15/92 10/15/92 10/15/92

METHOD BLANK SUMMARY

WU # 4432

LAB NAME : ENGINEERING-SCIENCE, INC. DATE ANALYZED :10/14/92

LAB SAMPLE ID:MWUG4921014

DATE EXTRACTED : NA

MATRIX :SOIL

INSTRUMENT ID: VGC-4

LAB	CLIENT	DATE
SAMPLE ID	Sample ID	ANALYZED
MWUG4921014 4432-2 4432-3 4432-4	METHOD BLANK HI-UW-4.5/-5.0/ HI-A-3/-4/ H式-UW-3/-3.5/	10/14/92 10/14/92 10/14/92 10/14/92

TOTAL RECOVERABLE PETROLEUM HYDROCARBONS DATA PACKAGE

ORGANIC ANALYTICAL REPORT

Work Order NO.: 4432

Parameter: TPH

Matrix: Soil

Unit: mg/Kg

Analytical

Method: 418.1

Date Extracted 10/27/92

QC Batch NO.: S92QCB026TPH

Date Analyzed: 10/29/92

		*======================================			
Sample ID:	С	lient ID:	Result	Reporting Limit	Percent Moisture
=========			********		
4432-01 4432-02 4432-03 4432-04 4432-05	HI- HI- 18 HZ-	VW-4'-4.5' VW-4.5'-5.0' A-3'-4' VW-3'-3.5' VW-4'-4.5'	22 15 ND 12	5 5 4 4	22.2 21.8 5.8 3.2
4432-05 4432-07 MSTPH921027	HI-	VW-4'-4.5' BKG-2.5'-3' HOD BLANK	13000 84 ND	4 5 4	6.1 11.4 NA

NA_ Not Analyzed ND_ Not Detected

ANALYST:

Man D

GROUP LEADER:

Repul

INORGANICS DATA PACKAGE

10/27/92

INORGANICS ANALYTICAL REPORT

Client: ES-Denver Work Order: 4432 Project: Matrix: Solid H\$→VW Client's ID: HI-BKG -3'-3.5' -2.5'-3' Sample Date: 10/03/92 10/03/92 10/04/92 % Moisture: Lab ID: 4432.04 4432.05 4432.06 Normal Parameter -----Results-----Report Method Units Date Limit Analyzed Alkalinity 90. ND ND SM 403(M) 50 mg/Kg CaCO3 11/02/92 3.2 Moisture 6.1 9.8 ASTM D2216 . 1 % by wt 10/22/92 рH 7.0 6.2 6.6 EPA 9045 NA pH Units

Samples for alkalinity analysis were extracted using 10mL water for each 1g sample. These water extracts were analyzed for alkalinity, and the results were calculated in the solid on a dry-weight basis.

NA- Not Applicable ND- Not Detected

GROUP LEADER:

5.8

6.0

nu

INORGANICS ANALYTICAL REPORT

Client: ES-Denver Work Order: 4432 Matrix: Project: AFCEE Solid Client's ID: HI-VW HI-VW HI-A -4'-4.5' -4.5'-5.0' -3'-4' 10/02/92 10/02/92 10/02/92 Sample Date: % Moisture: Lab ID: 4432.01 4432.02 4432.03 Normal -----Results-----Parameter Method Report Units Date Limit Analyzed 50 ND ND ND SM 403(M) mg/Kg CaCO3 11/02/92 Alkalinity 5.8 Moisture 22.2 21.8 ASTM D2216 . 1 Jw vd # 10/22/92

6.1

EFA 9045

NA

pH Units

10/27/92

Note: Samples for alkalinity analysis were extracted using 10mL water for each 1g sample. These water extracts were analyzed for alkalinity, and the results were calculated in the solid on a dry-weight basis.

NA- Not Applicable ND- Not Detected

ANALYST: Don Dleaton

GROUP LEADER:

INORGANIC OC SUMMARY - MS and MSD

Work Order:

4432

% Moisture:

NA

Lab ID Spk/Dup:

Alkalinity Moisture pH

Matrix:

Solid

QC Batch:

Blank Spk 4438.04 4454.01 452.42 451.82 453.40

Units: mg/Kg CaCO3 (Alk)

t by wt. (Hois) pH Units (pH)

	Date Analyzed	Results Unspiked		RPD	RPD QC	-Conc Ad	d e d -	Perc Recov	
Parameter	MS/Dup	Sample MS/Sample	KSD/Dup		Limit	NS	MSD	HS	KSD
Alkalinity Koisture pH	11/02/92 10/22/92 10/27/92	0.00 23000.00 8.33 5.93	7.52	0 11 2	20 29 26	23650.00	23650.00	97	ΥĪ

* or N = Outside QC Limit:

QC Limits for & Rec: 75 - 125

ANALYST: Don Sleator Date 11/11/92 REVIEWER: MUB Date 11/17/92. File: H1QCHSWH

INORGANICS ANALYTICAL REPORT

Client:

ES-Denver

Work Order:

4432

Project:

AFCEE

Matrix:

Solid

Client's ID:

Prep Blank

Sample Date:

% Moisture:

Lab ID:

Prep Blank

Parameter	Results	Method	Normal Report Limit	Units	Date Analyzed
Alkalinity	ND	SM 403(M)	50	mg/Kg CaCO3	11/02/92
Moisture	NA	ASTM D2216	.1	% by wt	10/22/92
pН	NA	EPA 9045	NA	pH Units	10/27/92

Note: Samples for alkalinity analysis were extracted using 10mL water for each 1g sample. These water extracts were analyzed for alkalinity, and the results were calculated in the solid on a dry-weight basis.

NA- Not Applicable ND- Not Detected

ANALYST:

Von Gleston

GROUP LEADER:

INORGANICS QC SUMMARY - LAB CONTROL SAMPLE

Work Order:

4432

% Moisture:

NA

Lab ID of LCS:

Alkalinity:

452.42 LCS

Matrix:

Solid

Units:

mg/Kg CaCO3

	Date Analyzed	LCS	Conc	% Rec	Advisory % Re	
Parameter	LCS	Result	Added	LCS	Low	High
Alkalinity	11/02/92	23000.00	23650.00	97	80	120

ANALYST: Don Bleaton Date 11/11/92 REVIEWER: 100B Date 1/17/92

METALS DATA PACKAGE

CASE NARRATIVE WORK ORDER NO. 4432 METALS

The serial dilution sample result for iron did not agree with the undiluted result within 10%, and the diluted sample result was greater than ten times the iron MDL. All iron results in this batch are therefore flagged with "E".

Client ID's were abridged by the laboratory to facilitate computer entry of analytical data. The following should be used as a reference:

CLIENT ID		ABRIDGED ID
H1-VW-4'-4.5'		H1VW-4
H1-VW-4.5'-5.0'		VW-4.5
H1-A-3'-4'	•	A-3-4
H2-VW-3'-3.5'		VW-3
H2-VW-4'-4.5'	gr - * · · · · · · · · · · · · · · · · · ·	H2VW-4
H1-BKG-2.5'-3'		BKG2.5

		INORGANIC	ANALYSES DATA S	SHEET	CLI	ENT SAM	PLE ID
						H1VW-4	1
.ab Name: E_S	BERKELEY_L	ABORATORY_	Contract: AF	CEE_ES-	-D		
ab Code: ESBI	L Ca	se No.: 44	32S SAS No.:		_ SDG	No.: A	-3-4_
fatrix (soil/	water): SOIL			Lab San	mple ID	: 4432.	01
evel (low/med	i): LOW_			Date Sa	ampled	: 10/02	/92
Solids:	_77.	8					
	oncentration	Units (ug	/L or mg/kg dry	weight	t): MG/	KG	
	CAS No.	 Analyte	 Concentration	ci ó	 M		
	7439-89-6	Iron	7980	E	P		
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omments:							
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		INORGANIC .	- ANALYSES DATA :	SH	EET	C.	LIENT SAMPLE ID
ah Nama R C	nanyaray r	**************************************	A	n a	nn na n		VW-4.5
			Contract: Al				
lab Code: ESBL	Ca	se No.: 44	32S SAS No.	• .		S	DG No.: A-3-4_
atrix (soil/w	ater): SOIL	_		L	ab Samp	le	ID: 4432.02
evel (low/med	LOW_	_		D	ate Sam	ple	d : 10/02/92
Solids:	_78.	2					
Co	ncentration	Units (ug	/L or mg/kg dry	У '	weight)	: M	G/KG
,	CAS No.	 Analyte	Concentration	l c	! Ω	М	
· · · · · ·	7439-89-6	Iron	6260	- -	Ē	P_	
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omments:							

		TNORGANIC	ANALYSES DATA S	SHEET	CI	LIENT SAMPLE ID
		INONOMMIC	Annii did bii i		1	A-3-4
Lab Name: E_S_	_BERKELEY_L	ABORATORY_	Contract: Al	FCEE_ES-	D _	N 3 3
Lab Code: ESBL	Ca	se No.: 44	32S SAS No.	:	SI	OG No.: A-3-4_
Matrix (soil/w	ater): SOIL			Lab Sam	ple 1	ID: 4432.03
Level (low/med): LOW_			Date Sa	mpled	d : 10/02/92
% Solids:	_94.	2				
Co	ncentration	Units (ug	/L or mg/kg dry	y weight): MC	e/KG
	CAS No.	Analyte	 Concentration	ici õ	М	· .
	7439-89-6	Iron	8630	_	P_	
Comments:						

		TNORGANTC	ANALYSES DATA	SHEET	CLIENT SAMPLE ID
	•	2101011111	DRIA	~	VW-3
ab Name: E_S_	_BERKELEY_L	ABORATORY_	Contract: A	FCEE_ES-D	
ab Code: ESBL	Ca:	se No.: 44	32S SAS No.	:	SDG No.: A-3-4_
atrix (soil/w	ater): SOIL	_		Lab Samp	le ID: 4432.04
evel (low/med): LOW	_		Date Sam	pled : 10/03/92
Solids:	_96.8	3			
Co	ncentration	Units (ug	/L or mg/kg dry	y weight)	: MG/KG
	I CAS No	1 2221111	 Concentration		
	l	l		l_l	<u> </u>
_ •	7439-89-6 	Iron	8160	_ <u>_</u> E	P_
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omments:					

INORGANIC ANALYSES DATA SHEET

CLIENT SAMPLE ID

Lab Name: E_S_	_BERKELEY_L	ABORATORY_	Contract: A	FCE	E_ES-I		H2VW-4	
Lab Code: ESBL							No.: A-3	-4_
Matrix (soil/w							: 4432.05	
Level (low/med): LOW	_		Da	te Sam	pled	: 10/03/9	2
% Solids:	_93.	9						
Co.	ncentration	Units (ug	/L or mg/kg dr	y w	eight)	: MG/	KG	
,	CAS No.	 Analyte	 Concentration	c	Q	М		
	7439-89-6	Iron	5460		E	P_		
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Comments:								-
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Inorganics Report CLIENT SAMPLE ID INORGANIC ANALYSES DATA SHEET BKG2.5 Lab Code: ESBL___ Case No.: 4432S SAS No.: ____ SDG No.: A-3-4_ Matrix (soil/water): SOIL_ Lab Sample ID: 4432.06____ Level (low/med): LOW___ Date Sampled : 10/04/92 Solids: _90.2 Concentration Units (ug/L or mg/kg dry weight): MG/KG ICAS No. | Analyte |Concentration|C| M 7439-89-6 | Iron 11600| | P_|

FORM I - IN

omments:

		TNODCANTO	ANALYSES DATA :	cuppm	CLIENT SAMPLE ID
		INORGANIC	ANALISES DAIA .	24661	
Lab Name: E_S_	_BERKELEY_L	ABORATORY_	Contract: Al	FCEE_ES-D	PBLANK
Lab Code: ESBL	Ca	se No.: 44	32S SAS No.	:	SDG No.: A-3-4_
Matrix (soil/w	ater): SOIL	_		Lab Samp	le ID: PBK 482.22
Level (low/med): LOW	-		Date Sam	oled : 11/04/92
% Solids:	100.0	2			
Co	ncentration	Units (ug	/L or mg/kg dry	y weight):	: MG/KG
	CAS No.	 Analyte	 Concentration		 M
	7439-89-6	 Iron	4.6	UE	P_
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comments:		•			

LABORATORY CONTROL SAMPLE (BLANK SPIKE)

Lab	Name:	E_SBERKI	ELEY_LABORATORY_	Contract:	AFCEE_ES-D					
Lab	Code:	ESBL	Case No.: 4432S	SAS No.:		SDG	No.:	A-3-4_		
ol.	id LCS	Source:	ESBL-LCSS					-		
Aqu	eous Lo	CS Source:								

	Aque	ous (ug/I		Solid (mg/kg)					
Analyte	True	Found	%R	True	Found	С	Limi	ts	% R
Iron				100.0 92.1 _ 80.0 12		120.0	0.0 _92.1		
	-					_ -			
						_ -			
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						_ -			
	.					- -			
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LABORATORY CONTROL SAMPLE (BLANK SPIKE)

ab Name:	E_SBERK	ELEY_LABORATORY_	Contract: AFCEE_ES-D					
Lab Code:	ESBL	Case No.: 4432S	SAS No.:		SDG	No.:	A-3-4_	
olid LCS	Source:	ESBL-LCSS						
queous L	CS Source:							

Analyte		eous (ug/L Found	 True	Solid Found (i (mg/kg) Lii	mits	& R
Iron			100.0	93.9	80.0	1120.0	93.9
	.						
	!! !!						
	.						

BLANK SPIKE DUPLICATE

CLIENT	SAMPLI	TD.

•							LCSS	 a
Lab Name: E	_sberkeley	_LABORATOR	Y_	Contract:	AFCEE_E	S-D		•
Lab Code: E	SBL	Case No.:	44325	SAS No).:		SDG No.:	A-3-4_
latrix (soi	1/water): S	oir_			Le	vel	(low/med):	_LOW
% Solids fo	r Sample: 1	00.0		8	Solids	for	Duplicate:	100.0

Concentration Units (ug/L or mg/kg as received):MG/KG

Analyte	Control Limit	-	C		RPD	Q	! M
Iron		92.0990		93.8830 _	1.9	<u> </u> _	P_
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ICP SERIAL DILUTION

EPA	SAMPLE	NO.
-----	--------	-----

8					 H1VW-4L
Lab	Name:	E_SBERKELEY	_LABORATORY_	Contract: AFCEE_ES-	
ab	Code:	ESBL	Case No.: 44325	S_ SAS No.:	SDG No.: A-3-4_

atrix (soil/water): SOIL_ Level (low/med): LOW___

Concentration Units: ug/L

1	1 11	Carial !	1 0 1	1 1	
ļ		Serial	8	!!	
. !	Initial Sample	Dilution	Differ-		
Analyte	Result (I) C	Result (S) C	ence	101	М
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Iron	66454.92	73844.92	11.1	IEIF	
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Method Detection Limits (Annually)

Lab Name:	E_SBERKE	LEY_LABORA	ATORY_	Contract	: AFCEE_ES	5-D	
ab Code:	ESBL	Case No.:	4432S_	SAS No.:		\$	SDG No.: A-3-4_
ICP ID Num	ber:	TJA_61_	М	Date:	08/31/92	2	
rlame AA I	D Number :			Matrix: 8	soir_		
Turnace AA	ID Number	:		(ug/L in	1.00g to	100r	ml digestate)
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	 Analyte	Wave- length (nm)		 	 MDL (ug/L)	М	
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Comments:							
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PREPARATION LOG

Lab Name: E_S_BERKELEY_LABORATORY_ Contract: AFCEE_ES-D

Lab Code: ESBL__ Case No.: _4432S_ SAS No.: ____ SDG No.:A-3-4_

ethod: P_

EPA Sample Preparation Weight Volume No. Date (gram) (mL) A-3-4				
No. Date (gram) (mL)		 Preparation	Weight	Volume
A-3-4	1		_	
BKG2.5	No.	Date	(gram)	((((((((((((((((((((
BKG2.5				100
H1VW-4				
H1VW-4				
H2VW-4	H1VW-4	1_11/04/92		
LCSS	1H2VW-4	1 11/04/92	1.24	
LCSSD		1 11/04/92	1.00	
MPA-9		11/04/92	1.00_	
MPB-9		11/04/92	1.04	100
PBLANK		11/04/92	1.21	100
VMP1-9 11/04/92 1.07 100 VMP2-9 11/04/92 1.13 100 VW-10 11/04/92 1.00 100 VW-3 11/04/92 1.03 100 VW-4.5 11/04/92 1.18 100				
VMP2-9 11/04/92 1.13 100 VW-10 11/04/92 1.00 100 VW-3 11/04/92 1.03 100 VW-4.5 11/04/92 1.18 100	IVMP1-9			100
VW-10 11/04/92 1.00 100 VW-3 11/04/92 1.03 100 VW-4.5 11/04/92 1.18 100	IVMP2-9			
VW-3		11/04/92	1.00	100
VW-4.511/04/921.18100		11/04/92	1.03_	100
	IVW-4.5	11/04/92	1.18	100
		11/04/92	1.01_	100
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ANALYSIS RUN LOG

_Lab Name: E_S__BERKELEY_LABORATORY_ Contract: AFCEE_ES-D

Lab Code: ESBL__ Case No.: 4432S_ SAS No.: ____ SDG No.:A-3-4_

Instrument ID Number: TJA 61 M_ Method: P_

Start Date: 11/09/92

End Date: 11/09/92

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1	ICSAB	1.00	1059	<u> </u>	<u> </u>	N .	<u> _ </u>	_	_	_	_	_	_	_	_	_	_!	!	_!	_!		_		_	_!	_!	!	-!	
	CRI	1.00	1104	l		_ _	 _	_	_	_	_	_1	_	_	_	_		_	_	_	_	_!	_	_	_!	_	-!	_!	
Ì	ZZZZZZ	1.00	1108	l		_1_	1_	_		_	_	_1	_	_	_	_	_	_	_	_	_	_	_!	_!	_!	_!	-!	_[
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i	LCSS	1.00	1117	l		_ X	_	_	_	_	_	_	_	_	_	_	_	_	_	_	_	_	_!		_!	_!	_	_!	
.	LCSSD	1.00	1122	 		_ X	1_	_	_	_	_		_		_	_	_	_	_	_			_	_!	_!	_!	-!	_!	_
	H1VW-4		1127	l		_ X	1_	_		_	_	_!	_	_!	_	_		_!	_	_	_	_	_		_!	_!	_!	_!	_
l	H1VW-4L_	1.00	1131	I		_ X	 _	_	_	_	_	_	_		_	_	_	_	_	_	_	_	_		_!	_	_!	-!	_
	CCV		1136	l		_ X	1_	_		_		_	_	_	_	_	_[_	_	_	_	_	!	_!		_!	_!	-	
_ i	CCB	1.00	1141	l		_ X	1_	_	<u> </u>	_	_	_	_	_	_	_	_			_	_	_	_	_	_	_!	_	_!	_
	VW-4.5	1.00	1145	l		_ X	1_	_	_	_	_	_	_	_	_	_!	_	_	_	_	_	_	_	_	_	_	_	-!	
	A-3-4	1.00	1150	l		_ X		_	_	_	_	_	_	_	_	_!	_!	!	_	_		_	_	_	_!	_!	_!	_!	_
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	H2VW-4	1.00	11159	l		_ X	1_	_	_	_	 _	_	_	_	_	_	_	i	_	_	_	_	_	_	_		_!	_	
l	BKG2.5	1.00	1204	l		_ X	1_	_	_	_	_	_	_	_	_	_	_	_	_	_	_	_	_	_	_	_!	_!	-!	_
ĺ	VW-7	1.00	1208	I		_ X	•	_	_	_	_	_1	_	_	_	_	_	_	_	_	_	_		_		_	_	_!	_
n İ	MPB-9	1.00	1213	I		_ X	,	_	_	_	_	_	_1		_	_	_	_	_		_	_	_	_	_	_!	_!	_!	_
i	MPA-9	1.00	1217	 		_ X	,	 _	_	 _	_	_	_	_	_	_	_	_	_	_	_	_	_	_	_	_!	_!	_!	_
•	ccv	1.00	1222	l		_ X	. —	 _	_	 _	_		_1	_	_	_			_	_	_	_	_	_	_!	_!	_!	_!	_
_	CCB	1.00	1227	I		_ X	. —	_	_	 _	_	_	_	_	_	_	_	_		_	_	_	_	_	_!	_!	_[_!	_
İ	VW-10	1.00	1231	l		_ X	. —	_	_	1_	_	_	_	_	_	_1	!		_	_		_			!		_[_!	_
	VMP2-9		1236	l		_ X	. —	 _	_	1_	_		_1	_	_	_	_		_	<u> </u> _	_	_	_	_	_[_	_!	_!	_!	-
İ	VMP1-9	1.00	1241	 		_ X		1_	_	1_	_	_	_	_	_	_		_	_	_	_	_	!	_		_!	_!	_!	_
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ILM02.1

ANALYSIS RUN LOG

_Lab Name: E_S__BERKELEY_LABORATORY_ Contract: AFCEE_ES-D

Lab Code: ESBL__ Case No.: 4432S_ SAS No.: ____ SDG No.: A-3-4_

Instrument ID Number: TJA 61 M_ Method: P_

Start Date: 11/09/92

End Date: 11/09/92

													Αn	al	yt	es											1
EPA Sample No.	D/F	Time	% R	F																 -							_!
CRI	1.00	1254 1259		_ X				_ _	_		_ _	_		_ _	_ _	_ .	_! _!	_	_ _	_ _!	_	_ _!	_i _!	_ _	_ . _ .	_ - - -	- i
CCB	1.00	1304		_ _ _ X	_	-	_ _	_	_	_	_ _	_ _	_ _	_ _	_ _	_ _	_ _	_ _	_ _			! -	! !	- -	- - -	- - - -	_ i
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ILMO2.1

TOTAL PHOSPHORUS

TOTAL KJELDAHL NITROGEN

SOIL CLASSIFICATION

DATA PACKAGE



Engineering Science, Inc. 600 Bancroft Way Berkeley, CA 94710 Attention: Tom Paulson Client Project ID: W.O. #4432

Sample Descript: Soil

Analysis for: Percent Solids
First Sample #: 210-0637

Sampled: Received: 9/2-4/92 Oct 7, 1992

Analyzed:

Oct 19, 1992

Reported: Oct 20, 1992

LABORATORY ANALYSIS FOR:

Percent Solids

Sample Number	Sample Description	Detection Limit %	Sample Result %
210-0637	H1-W-4'-4.5'	10	83
210-0638	H1-VW-4.5'-5.0'	10	84
210-0639	H1-A-3'-4'	10	94
210-0640	H2-VW-3'-3.5'	10	97
210-0641	H2-VW-4'-4.5'	10	92
210-0642	H1-BKG-2.5'-3'	10	92

Analytes reported as N.D. were not present above the stated limit of detection.

SEQUOIA ANALYTICAL

Tod Granicher Project Manager THIS REPORT HAS BEEN

APPROVED AND REVIEWED BY

ESBL PROJECT MANAGER

DAI

600 Bancroft Way Berkeley, CA 94710

Attention: Tom Paulson

Client Project ID: Sample Descript:

: W.O. #4432 t: Soil

Total Phosphorous

Analysis for: Total Pho First Sample #: 210-0637 Sampled:

9/2-4/92

Received: Analyzed: Oct 7, 1992 Oct 19, 1992

Reported: Oct 20, 1992

LABORATORY ANALYSIS FOR:

Total Phosphorous

Sample Number	Sample Description	Detection Limit mg/kg	Sample Result mg/kg	
210-0637	H1-VW-4'-4.5'	10	370	
210-0638	H1-VW-4.5'-5.0'	10	290	
210-0639	H1-A-3'-4'	10	300	
210-0640	H2-VW-3'-3.5'	10	450	
210-0641	H2-VW-4'-4.5'	10	600	
210-0642	H1-BKG-2.5'-3'	10	460	
•	Method Blank	1.0	N.D.	

Analytes reported as N.D. were not present above the stated limit of detection.

SEQUOIA ANALYTICAL

sled

Tod Granicher Project Manager Please Note:

Analysis results reported on a dry-weight basis.

2100637.ENG <8>

Engineering Science, Inc. 600 Bancroft Way

Attention: Tom Paulson

Client Project ID:

W.O. #4432

Sampled:

9/2-4/92

Berkeley, CA 94710

Sample Descript: Analysis for:

Soil Total Kjeldahl Nitrogen Received: Analyzed:

Oct 7, 1992 Oct 15, 1992

First Sample #:

210-0637

Reported:

Oct 20, 1992

LABORATORY ANALYSIS FOR:

Total Kjeldahl Nitrogen

Sample Number	Sample Description	Detection Limit mg/kg	Sample Result mg/kg
210-0637	H1-VW-4'-4.5'	20	1,100
210-0638	H1-VW-4.5'-5.0'	20	730
210-0639	H1-A-3'-4'	20	70
210-0640	H2-VW-3'-3.5'	20	66
210-0641	H2-VW-4'-4.5'	20	53
210-0642	H1-BKG-2.5'-3'	20	91
-	Method Blank	0.10	N.D.

Analytes reported as N.D. were not present above the stated limit of detection.

SEQUOIA ANALYTICAL

Joe !!

Tod Granicher Project Manager Please Note:

Analysis results reported on a dry-weight basis.



600 Bancroft Way

Berkeley, CA 94710

Attention: Tom Paulson

Client Project ID: W.O. #4432

QC Sample Group: 210-0637-42

Reported: Oct 20, 1992

QUALITY CONTROL DATA REPORT

ANALYTE		Total Kjeldahl					
L	Total Phosphorous	Nitrogen					
Method: Analyst:	EPA365.3 K. Follett	EPA351.4 G. Kern		•	• • • •		
Reporting Units: Date Analyzed: QC Sample #:	mg/kg Oct 19, 1992 210-0642	mg/kg Oct 15, 1992 210-0642			•	•	
ao odmpie #.	210-00-12	210-00-12	•			••	
Sample Conc.:	420	84	. 1.1 m - 1. v m	e e e e	•• . • • . • •		• • • • • • • • • • • • • • • • • • • •
Spike Conc. Added:	100	4000					
Conc. Matrix Spike:	560	4000					
Matrix Spike % Recovery:	140	98				. · · .	
Conc. Matrix Spike Dup.:	530	3800					1. 1.
Matrix Spike Duplicate % Recovery:	110	93					
Relative % Difference:	5.6	5.1					

SEQUOIA ANALYTICAL

Joell

Tod Granicher Project Manager

% Recovery:	Conc. of M.S Conc. of Sample	x 100	
	Spike Conc. Added	•	
Relative % Difference:	Conc. of M.S Conc. of M.S.D.	x 100	
	(Conc. of M.S. + Conc. of M.S.D.) / 2		

2100637.ENG <10>



600 Bancroft Way Berkeley, CA 94710

Attention: Tom Paulson

Client Project ID: Sample Descript:

Lab Number:

W.O. #4432 Soil, H1-VW-4'-4.5'

Method of Analysis: ASTM D422-63

210-0637

Sampled: Received:

Sep 2, 1992 Oct 7, 1992

Analyzed: Reported:

Oct 13, 1992 Oct 20, 1992

PARTICLE SIZE DISTRIBUTION BY SIEVE AND HYDROMETER

SIEVE TEST

(A) TOTAL WEIGHT OF SAMPLE:

(B) WEIGHT RETAINED IN NO. 10 SIEVE:

(C) % PASSING NO. 10 SIEVE:

218.37g 33.38g 84.71

SIEVE TEST FOR WEIGHT RETAINED IN NO. 10 SIEVE

IDEAL PAN = 0.0 IDEAL TOTAL = (B)

	WEIGHT		CUMULATIVE	CUMULATIVE
SIEVE SIZE	RETAINED, g	% RETAINED	% RETAINED	% PASSING
1½ in.	0.0	0.0	0.0	100
3/8 in.	7.00	3.21	3.21	96.79
No. 4	10.54	4.83	8.04	91.96
No. 10	15.84	7.25	15.29	84.71
PAN	0.0	ĺ		•
TOTAL	33.38			

HYDROMETER TEST

ELAPSED TIME	TEMP.	HYDROMETER	CORRECTED		PARTICLE
(T)	°C	READING (H)	READING (R)	(L)	DIAM. (S)
2	21	20	16	13.7	0.035
5	21	17	13	14.2	0.023
10	21	15	11	14.5	0.016
15	21	15	11	14.5	0.013
25	21	13	9	14.8	0.010
40	21	12	8	15.0	0.0083
60	21	11	7	15.2	0.0068
90	21.	11	7	15.2	0.0055
120	21	10	6	15.3	0.0048
1440	21	8	4	15.6	0.0014

% SUSPENDED
(P)
21
17
15
15
12
11
9.3
9.3
8.0
5.3

& CHICDENIDED

WEIGHT OF SOIL USED IN HYDROMETER TEST (D): HYGROSCOPIC MOISTURE CORRECTION FACTOR (G):

SPECIFIC GRAVITY (ASSUMED):

DISPERSING AGENT CORRECTION FACTOR (E):

MENISCUS CORRECTION FACTOR (F):

TEMP./SPEC. GRAVITY DEPENDANT CONSTANT (K):

	65g	l
	0.983	۱
	2.65	I
1	3	١
	1	١
	0.01348	١

FORMULAS:

R = H - E - F

S = K[SQRT(L/T)]P = (R/W) 100

 $W = (J \cdot 100) / C$

 $J = D \cdot G$

SEQUOIA ANALYTICAL

Tod Granicher Project Manager

2100637.ENG <1>

600 Bancroft Way

Client Project ID:

W.O. #4432

Sampled:

Sep 2, 1992

Berkeley, CA 94710

Sample Descript: Soil, H1-VW-4.5'-5.0' Method of Analysis: ASTM D422-63

Received: Analyzed:

Oct 7, 1992 Oct 13, 1992

Attention: Tom Paulson

Lab Number:

210-0638

Reported:

Oct 20, 1992

PARTICLE SIZE DISTRIBUTION BY SIEVE AND HYDROMETER

SIEVE TEST

(A) TOTAL WEIGHT OF SAMPLE:

(B) WEIGHT RETAINED IN NO. 10 SIEVE:

(C) % PASSING NO. 10 SIEVE:

255.17g 4.75a 98.14

SIEVE TEST FOR WEIGHT RETAINED IN NO. 10 SIEVE

> IDEAL PAN = 0.0 IDEALTOTAL = (B)

	WEIGHT		CUMULATIVE	CUMULATIVE
SIEVE SIZE	RETAINED, g	% RETAINED	% RETAINED	% PASSING
1½ in.	0.0	0.0	6.0	100
3/8 in.	0.0	0.0	0.0	100
Nc. 4	1.48	0.58	0.58	99.42
No. 10	3.27	1.28	1.86	98.14
				
PAN	0.0		<u> </u>	

4.75 TOTAL

HYDROMETER TEST

ELAPSED TIME	TEMP.	HYDROMETER	CORRECTED		PARTICLE
(T)	°C	READING (H)	READING (R)	(L)	DIAM. (S)
2	. 21	12	8	15.0	0.037
5	21	10	6	∷15.3	0.024
10	21	9	5	15.5	0,017
15	21	8	4	15.6	0.014
25	21	8	4	15.6	0.011
40	21	8	4	15.6	0.0084
60	21	7	3	15.8	0.0069
90	21	7	3	15.8	0.0056
120	21	7	3	15.8	0.0049
1440	21	7	3	15.8	0.0014

	% SUSPENDED
	(P)
	12
	9.2 7.6
	6.1
	6.1
	6.1
İ	4.6
	4.6
1	4.6
	4.6

WEIGHT OF SOIL USED IN HYDROMETER TEST (D): HYGROSCOPIC MOISTURE CORRECTION FACTOR (G): SPECIFIC GRAVITY (ASSUMED): DISPERSING AGENT CORRECTION FACTOR (E):

MENISCUS CORRECTION FACTOR (F):

TEMP./SPEC. GRAVITY DEPENDANT CONSTANT (K):

65g FORMULAS: R = H - E - F0.983

S = K[SQRT(L/T)]2.65 3 P = (R/W) 100

 $W = (J \cdot 100) / C$

 $J = D \cdot G$

1 0.01348

SEQUOIA ANALYTICAL

Tools

Tod Granicher Project Manager

2100637.ENG <2>



600 Bancroft Way

Client Project ID: Sample Descript: Soil, H1-A-3'-4'

W.O. #4432

Sampled:

Sep 2, 1992 Oct 7, 1992

Berkeley, CA 94710

Method of Analysis: ASTM D422-63

Received: Analyzed:

Oct 13, 1992

Attention: Tom Paulson

Lab Number:

210-0639

Reported:

Oct 20, 1992

PARTICLE SIZE DISTRIBUTION BY SIEVE AND HYDROMETER

SIEVE TEST

(A) TOTAL WEIGHT OF SAMPLE:

(B) WEIGHT RETAINED IN NO. 10 SIEVE:

(C) % PASSING NO. 10 SIEVE:

256.56a 99.81g

SIEVE TEST FOR WEIGHT RETAINED IN NO. 10 SIEVE

IDEAL PAN = 0.0 IDEAL TOTAL = (B)

	WEIGHT		CUMULATIVE	CUMULATIVE
SIEVE SIZE	RETAINED, g	% RETAINED	% RETAINED	% PASSING
1½ in.	0.0	0.0	0.0	100
3/8 in.	41.21	16.06	16.06	83.94
"No. 4	26.13	10.18 ***	26.24	73.76
No. 10	32.47	12.66	36.90	61.10
DAN	0.0	i		

PAN 0.0TOTAL 99.81

HYDROMETER TEST

ELAPSED TIME	TEMP.	HYDROMETER	CORRECTED		PARTICLE
(T)	°C	READING (H)	READING (R)	(L)	DIAM. (S)
- 2	21	14	-10	14.7	0.037
5	21	12	8	15.0	0.023
10	21	11	7	15.2	0.017
15	21	10	6	15.3	0.014
25	-21	9	5	15.5	0.011
40	21	9	5	15.5	0.0084
60	21	7	3	15.8	0.0069
90.	21	7	3	15.8	0.0056
120	21	7	3	15.8	0.0049
1440	21	7	3	15.8	0.0014

2 OUSPENDED
(P)
9.4
7.6
6.5
5.7
4.7
4.7 2.8 2.8 2.8
2.8
2.8
2.8
2.8

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WEIGHT OF SOIL USED IN HYDROMETER TEST (D): HYGROSCOPIC MOISTURE CORRECTION FACTOR (G): SPECIFIC GRAVITY (ASSUMED):

DISPERSING AGENT CORRECTION FACTOR (E):

MENISCUS CORRECTION FACTOR (F):

TEMP./SPEC. GRAVITY DEPENDANT CONSTANT (K):

65g FORMULAS: 0.995 R = H - E - F2.65

3

1

0.01348

S = K[SQRT(L/T)]P = (R/W) 100 $W = (J \cdot 100) / C$

 $J = D \cdot G$

SEQUOIA ANALYTICAL

Tod Granicher Project Manager

600 Bancroft Way

Berkeley, CA 94710

Client Project ID:

W.O. #4432

Sampled:

Sep 4, 1992

Sample Descript: Soil, H1-BKG-2.5'-3' Method of Analysis: ASTM D422-63

Received: Analyzed:

Oct 7, 1992 Oct 14, 1992

Attention: Tom Paulson

Lab Number:

210-0642

Reported:

Oct 20, 1992

PARTICLE SIZE DISTRIBUTION BY SIEVE AND HYDROMETER

SIEVE TEST

(A) TOTAL WEIGHT OF SAMPLE:

(B) WEIGHT RETAINED IN NO. 10 SIEVE:

(C) % PASSING NO. 10 SIEVE:

280.69a 39.34g 85.98

SIEVE TEST FOR WEIGHT RETAINED IN NO. 10 SIEVE

IDEAL PAN = 0.0 IDEAL TOTAL = (B)

	WEIGHT		CUMULATIVE	CUMULATIVE
SIEVE SIZE	RETAINED, g	% RETAINED	% RETAINED	% PASSING
1½ in.	0.0	0.0	0.0	100
3/8 in.	24.76	8.62	8.82	91.18
No. 4	7.50	2.67	11.49	88.51
No. 10	7.08	2.52	14.01	85.99
		•		
PAN	1 00	<u> </u>	<u> </u>	

TOTAL 39.34

HYDROMETER TEST

ELAPSED TIME	TEMP.	HYDROMETER	CORRECTED		PARTICLE
(T)	°C	READING (H)	READING (R)	(L)	DIAM. (S)
2	21	- 26	22	12.7	0.034
5	21	24	20	13.0	0.022
10	21	24	20	13.0	0.015
15	21	22	18	13.3	0.013
25	21	19	15	13.8	0.010
40	21	19	15	13.8	0.0079
60	21	15	11	14.5	0.0066
90	21	15	11	14.5	0.0054
120	21	12	8	15.5	0.0048
1440	21	9	5	15.5	0.0014

(P)
30
27
22
24
20
20
15
15
11
6.7

% SUSPENDED

WEIGHT OF SOIL USED IN HYDROMETER TEST (D): HYGROSCOPIC MOISTURE CORRECTION FACTOR (G):

SPECIFIC GRAVITY (ASSUMED):

DISPERSING AGENT CORRECTION FACTOR (E):

MENISCUS CORRECTION FACTOR (F):

TEMP./SPEC. GRAVITY DEPENDANT CONSTANT (K):

	65g
	0.0984
	2.65
	3
Ì	1
	0.01348

FORMULAS:

R = H-E-F

S = K[SQRT(L/T)]P = (R/W)100

 $W = (J \cdot 100) / C$

 $J = D \cdot G$

SEQUOIA ANALYTICAL

Tod Granicher Project Manager

2100637.ENG <6>

600 Bancroft Way Berkeley, CA 94710 Attention: Tom Paulson

Client Project ID: Sample Descript:

Lab Number:

W.O. #4432 Soil, H2-VW-3'-3.5'

Method of Analysis: ASTM D422-63 210-0640

Sampled: Received: Sep 3, 1992 Oct 7, 1992

Analyzed: Reported:

Oct 13, 1992 Oct 20, 1992

PARTICLE SIZE DISTRIBUTION BY SIEVE AND HYDROMETER

SIEVE TEST

(A) TOTAL WEIGHT OF SAMPLE:

(B) WEIGHT RETAINED IN NO. 10 SIEVE:

(C) % PASSING NO. 10 SIEVE:

203.97a 33.37g 88.64

TOTAL

33.37

SIEVE TEST FOR WEIGHT RETAINED IN NO. 10 SIEVE

IDEAL PAN = 0.0 IDEAL TOTAL = (B)

	WEIGHT		CUMULATIVE	CUMULATIVE
SIEVE SIZE	RETAINED, g	% RETAINED	% RETAINED	% PASSING
1½ in.	0.0	. 0.0	0.0	100
3/8 in.	4.04	1.96	1.98	98.02
No. 4	15.8 i	7.75	9.73	90.27
No. 10	13.52	6.63	16.36	83.64
	<u> </u>	<u> </u>		
PAN .	1 00			

HYDROMETER TEST

ELAPSED TIME	TEMP.	HYDROMETER	CORRECTED		PARTICLE
(T)	°C	READING (H)	READING (R)	(L)	DIAM. (S)
2	21	9	5	15.5	0.038
5	21	9	5	15.5	0.024
10	21	8	4	15.6	0.017
15	21	. 7	3	15.8	0.014
25	21	7	3	15.8	0.011
40	21	7	3	15.8	0.0085
60	21	6	2	16.0	0.0070
90	21	5	1	16.1	0.0057
120	21	5	1	16.1	0.0049
1440	21	5	1	16.1	0.0014

% SUSPENDED (P) 5.5 **G.5** 5.2 3.9 3.9 3.9 2.6 1.3 1.3 1.3

WEIGHT OF SOIL USED IN HYDROMETER TEST (D): HYGROSCOPIC MOISTURE CORRECTION FACTOR (G): SPECIFIC GRAVITY (ASSUMED):

DISPERSING AGENT CORRECTION FACTOR (E): MENISCUS CORRECTION FACTOR (F):

TEMP./SPEC. GRAVITY DEPENDANT CONSTANT (K):

65g 0.983 2.65 3 1 0.01348

FORMULAS:

R = H-E-F S = K[SQRT(L/T)]

P = (R/W)100

 $W = (J \cdot 100) / C$

 $J = D \cdot G$

SEQUOIA ANALYTICAL

Tod Granicher **Project Manager**

600 Bancroft Way

Berkeley, CA 94710 Attention: Tom Paulson

Client Project ID: Sample Descript:

Lab Number:

W.O. #4432 Soil, H2-VW-4'-4.5'

Method of Analysis: ASTM D422-63 210-0641

Sep 3, 1992 Oct 7, 1992 Oct 14, 1992

Received: Analyzed: Reported:

Sampled:

Oct 20, 1992

PARTICLE SIZE DISTRIBUTION BY SIEVE AND HYDROMETER

SIEVE TEST

(A) TOTAL WEIGHT OF SAMPLE:

(B) WEIGHT RETAINED IN NO. 10 SIEVE:

(C) % PASSING NO. 10 SIEVE:

228.60a 7.04q 96.92

SIEVE TEST FOR WEIGHT RETAINED IN NO. 10 SIEVE

IDEAL PAN = 0.0 IDEALTOTAL = (B)

	WEIGHT		CUMULATIVE	CUMULATIVE
SIEVE SIZE	RETAINED, g	% RETAINED	% RETAINED	% PASSING
1½ in.	0.0	0.0	0.0	100
3/8 in.	0.0	0.0	0.0	100
No. 4	2.50g	1.09	1.09	98.91
No. 10	4.54g	1.99	3.08	95.92
	İ			
PAN	0.0		7	

TOTAL 7.04

HYDROMETER TEST

ELAPSED TIME	TEMP.	HYDROMETER	CORRECTED		PARTICLE
(T)	°C	READING (H)	READING (R)	(L)	DIAM. (S)
2	21	7	3	15.8	0.038
5	21	7	3	15.8	0.024
10	21	7	3	15.8	0.017
15	21	6	2	16.0	0.014
25	-21	5	i i	16.1	-0.011
40	. 21	5	1 1	16.1	0.0086
60	21	5	1	16.1	0.0070
90	21	5	1	16.1	0.0057
120	21	5	1	16.1	0.0049
1440	21	5	i	16.1	0.0014

% SUSPENDED
(P)
4.5
4.5
4.5
3.0
1.5
1.5
1.5
1.5
1.5
1.5

WEIGHT OF SOIL USED IN HYDROMETER TEST (D): HYGROSCOPIC MOISTURE CORRECTION FACTOR (G):

SPECIFIC GRAVITY (ASSUMED):

DISPERSING AGENT CORRECTION FACTOR (E):

MENISCUS CORRECTION FACTOR (F):

TEMP./SPEC. GRAVITY DEPENDANT CONSTANT (K):

65g	FO
0.995	
2.65	
3	
1	
0.01348	

RMULAS:

R = H - E - FS = K[SQRT(L/T)]

P = (R/W)100

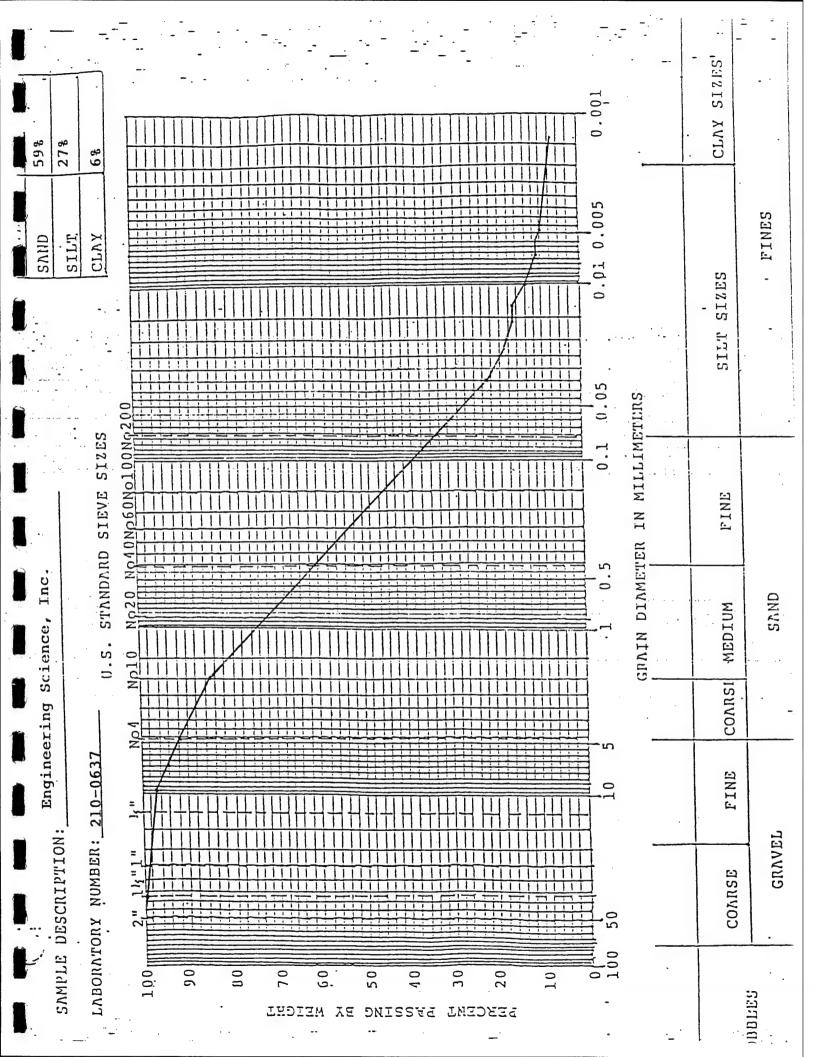
 $W = (J \cdot 100) / C$

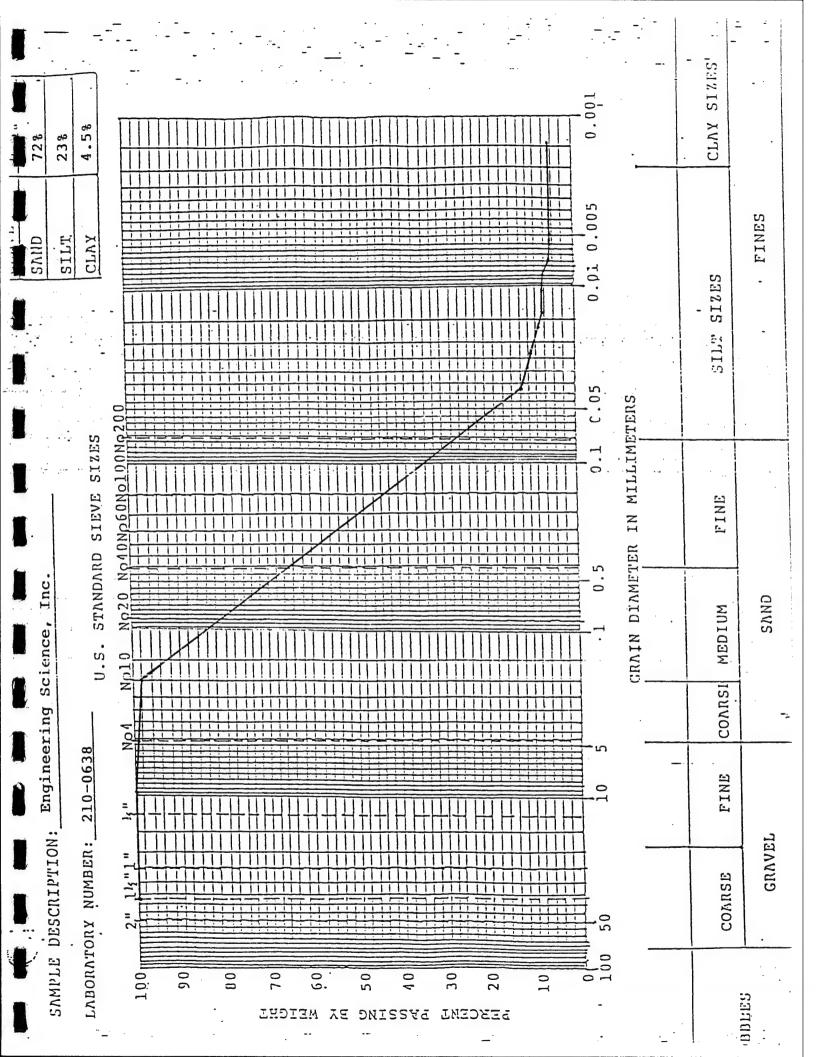
 $J = D \cdot G$

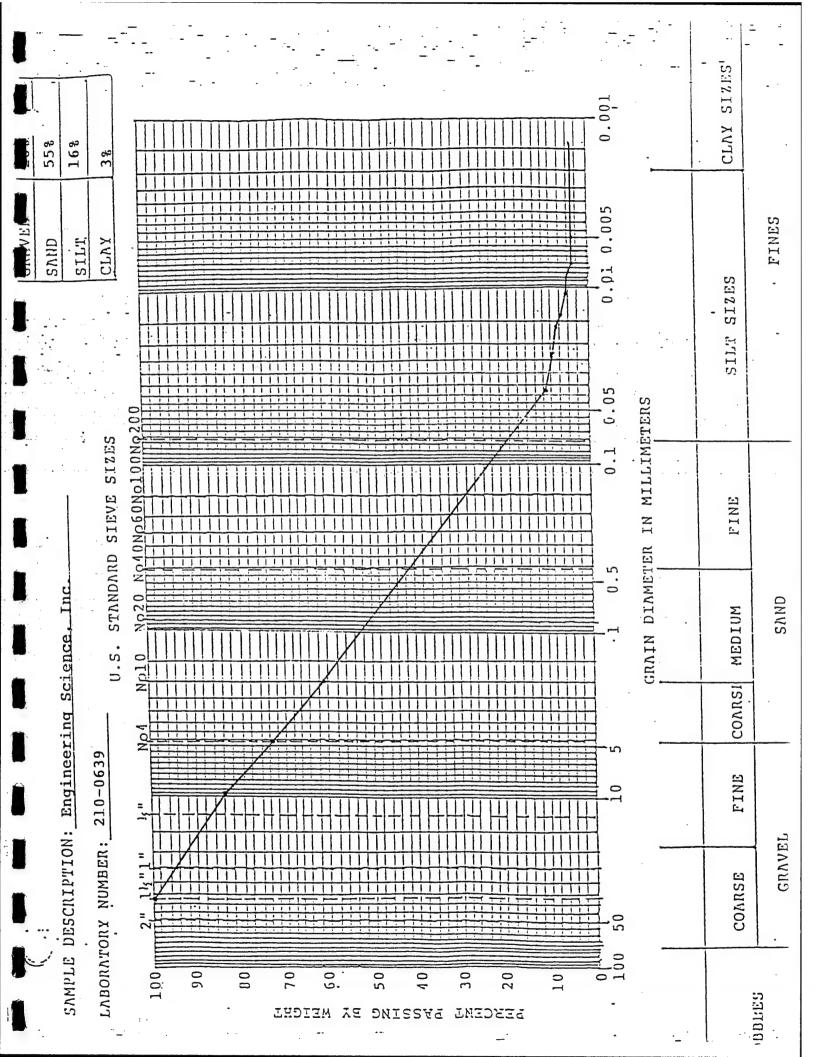
SEQUOIA ANALYTICAL

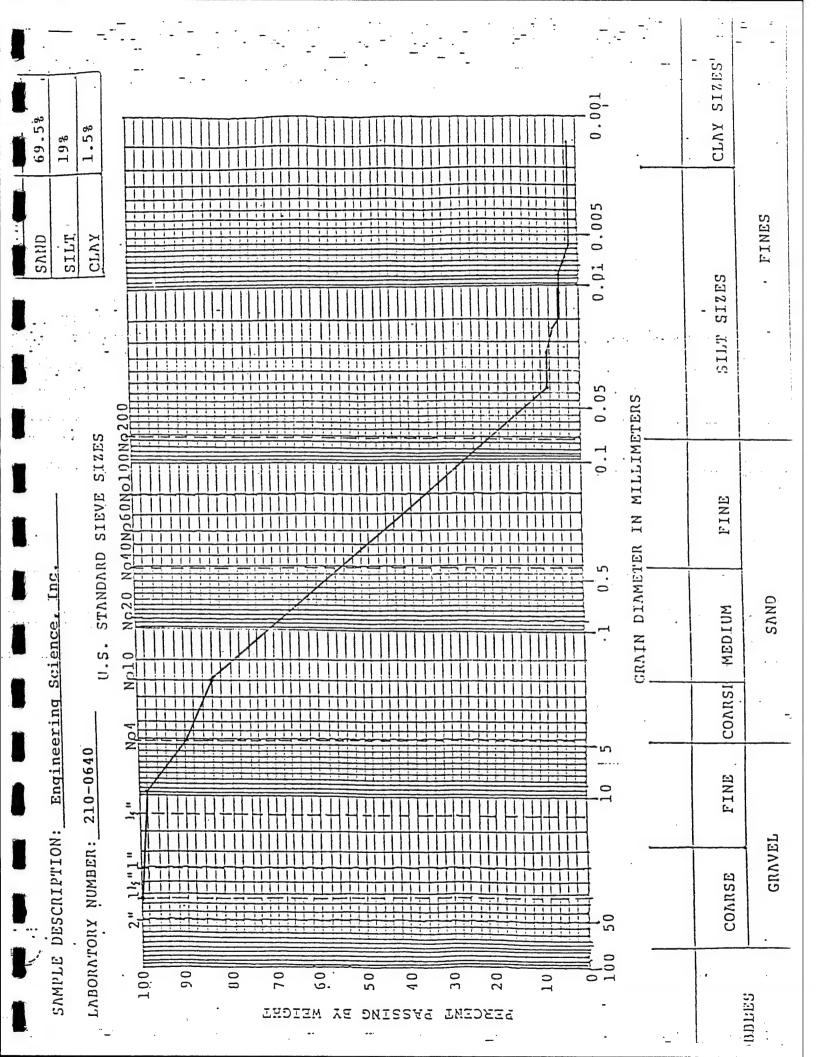
Steel

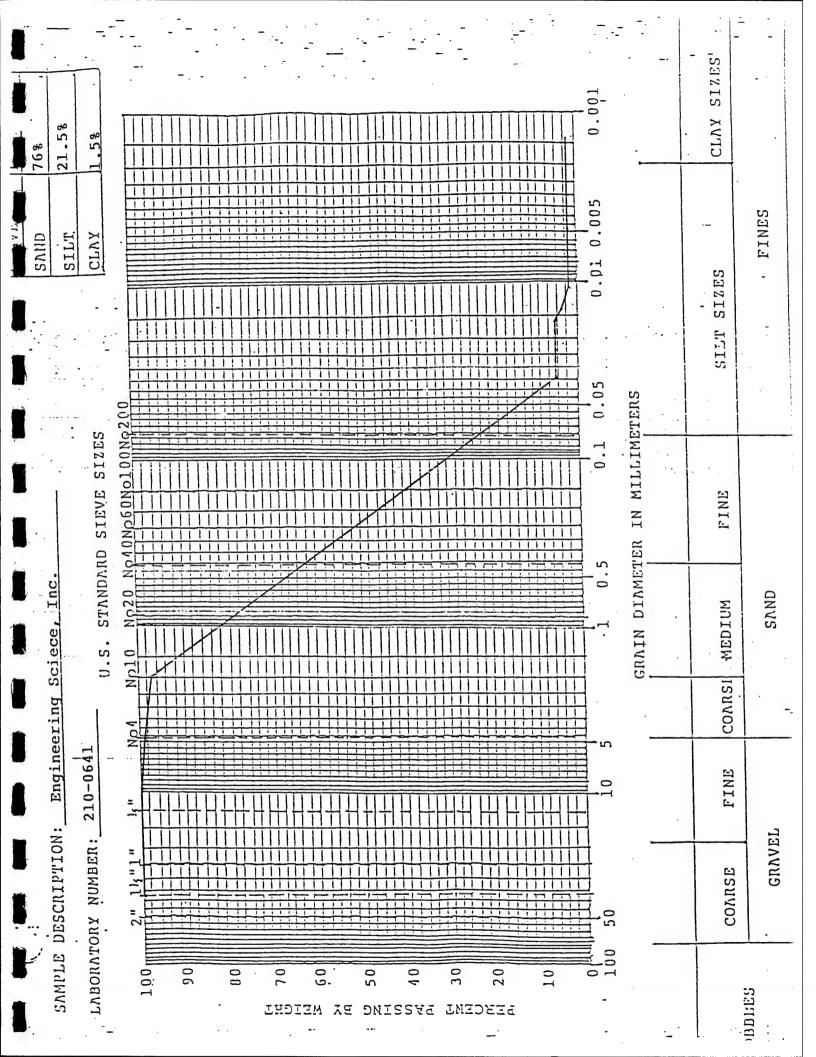
Tod Granicher Project Manager

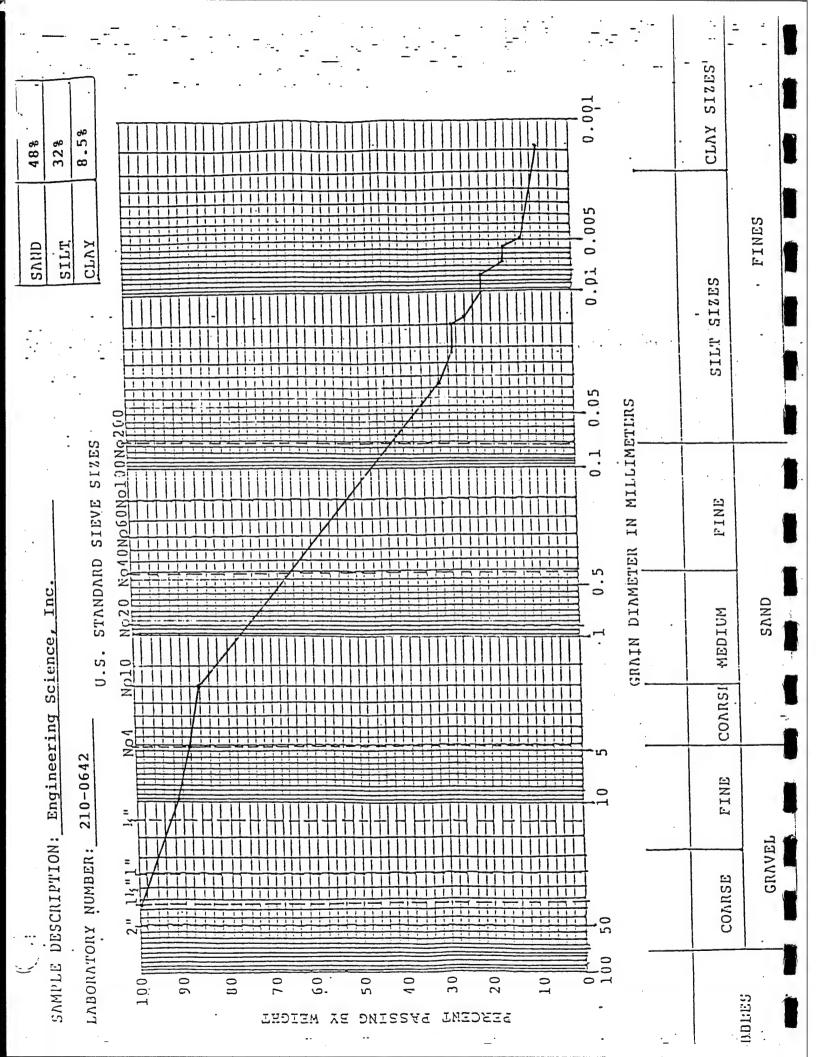












ENGINEERING-ECTENCE

CHAIN OF CUSTODY RECORD FOR WATER BAMPLES

ំ ខា បាល	р 110.	PROJECT NAME/LOCATION		PREBERVATIVES	virves required	
·		4432		ارمدد		To: STOUGHA LAB.
FIELD C	CONTACT:			A ANALYS	ев педиливр	
BAMPLER	BAMPLERB NAMES	1 & BIGNATURES		0)5.		
			314H			
,			45346 Y.	(NY 31		
DATE	ттие	FIELD SAMPLE IDENTIFIER	741	12/5		пенликв
9/2/92	15.60	HI-VW-4'-45' (443).018	/	7	3100637	Word nosult as Any
	1500	141.VW-4,51-5,0' (4432.02B)	\ \ '	7	88	BASIS CULE MIX/13. FO
	1,500	141-4-3-4' (4432038)	/	>	£ 29	Hoponthub ant to.
1/3/6,2	1200	42.VW- 3- 3.5 (4432,044)V	\ \ \ \	7	922	Urpart meties Blow
	*	42-1W-4-45'(4422,058)	\	7	5,00	45/450 . (10 day next
4/91		H1-BKG- 2,5-3 (443). W.B.	1/0	7	J C640	TAT.) NOPONT AND
				٠		To Ton Payleon.
FIELD C	UBTODY	cubrody refinguibled by:		El cire on	(), DATE:	: 10/6/92 TIME: 16:40
ингерер	VIN:	AIRBILL #	NO	RECEIPT	CUBTODY BEALB?	, TEMP:
RECEIVED		FOR LABORATORY DY:	les.		DATE:	: 10 / 6/92 TIME: 4:00 Am
20 mg	1180	106/90	9-	1	19-20	1125

Baffelle

Columbus Laboratories

アローショウン JOB No. Engineering-Science 200 300

CHAIN OF CUSTCDY RECORD

Form No.

90

402 glASS/1602 G1ASS BRASS TUBE BRASS TUBE BRASS FUBE SLASS TUBE 1602, 6/455 402 GIASS BLKS TUBE BASS TOBE 1602, 61455 16-2 6/45 6/455 6/15/ 1602 6/ASS Remarks 402 GlASS 402, GIASS 402 6/ASS 20% 7091 Received by: Received by: (Signature) (Signature) RESULTS TO! Containers ło Митрег Container No. JEFF K: HEL SOS KING AUE, Date/Time Date/Time Remarks SEND Al min Park SAMPLE TYPE (V) Relinquished by: (Signature) 26/19/01 Relinquished by: (Signature) X ¥ يح بح 10/06/2 0930 $\frac{\times}{\lambda}$ Date/Time × × ン × × کر × × × Received for Laboratory by ス > Received by: (Signature) HI-BK6-3'-35' HI-A-3583-4 141-BK6-2,5'-3' -BK6 -2,5'-3' - WW- 4.5 - 5.0 Received by: 42-VW-3'-3.5' 42-UW-3-3,51 (Signature) Signa; http: 13-14-4121 13-10-4-6.5 54-, t-M1-42 -VW-4.5 5-4-14-M1-1H 11-VW - 4-45 SAMPLE I.D. HI-A-3-4' -VW-4,5 HI-A-3-4 -VW-4.5 A FB 104 CCT 92 2200 Date/Time Date/Time Date/Time Project Title HANSGER アング Relinquished by: (Signature) TIME Relinquished by: (Signature) Relinquished by: Signature) leuly M SAMPLERS: (Signature) mx Hearing 0490-87hhS 03 OCT 92 20x792 0304792 020CT 92 04 KT 92 34 OCT 92 02 01 92 0306792 03007 92 0400792 32 OCT 92 03 WT 92 130ct 92 02067 92 52 0CT 92 03 OT 92 52067 92 DATE Proj. No.

APPENDIX C BUILDING 1639 SOIL GAS PERMEABILITY DATA

Table C-1. Results of Soil Gas Permeability Test at Monitoring Point H1-MPA

0 0.05 0.02 0.03 0.03 0.03 0.03 0.03 0.02 0.02	rressure ("H ₂ O) by Deptn		Pressure ("H	Pressure ("H ₂ O) by Depth
0 0.05 0.02 0.03 0.03 0.03 0.03 0.02 0.02 0.015	5.0′	Time (min)	2.5′	5.0′
0.05 0.02 0.03 0.03 0.03 0.03 0.02 0.02 0.015	0.01	20	0.025	0.32
0.02 0.03 0.033 0.035 0.03 0.02 0.02 0.015	0.17	23	0.025	0.33
0.03 0.033 0.035 0.03 0.02 0.02 0.015	0.21	26	0.025	0.34
0.03 0.033 0.03 0.02 0.02 0.015	0.22	29	0.03	0.35
0.033 0.35 0.03 0.02 0.015 0.015	0.225	32	0.025	0.35
0.35 0.03 0.02 0.015 0.015	0.235	35	0.03	0.35
0.02 0.02 0.015	0.25	38	0.025	0.35
0.02 0.015 0.015	0.27	41	0.025	0.35
0.02	0.28	44	0.02	0.35
0.015	0.30	47	0.02	0.35
0.01	0.30	50	0.03	0.35
	0.30	60	0.02	0.35
14 0.02 0.30	0.30	70	0.02	0.36
16 0.02 0.30	0.30	80	0.01	0.37
18 0.03 0.32	0.32	06	0.02	0.36

Table C-2. Results of Soil Gas Permeability Test at Monitoring Point H1-MPB

	Pressure ("H	Pressure ("H ₂ O) by Depth		Pressure ("E	Pressure ("H ₂ O) by Depth
Time (min)	2.5′	5.0′	Time (min)	2.5′	5.0′
1	0.045	0.01	12	0.080	0.015
2	90:0	0.02	50	0.115	0.025
3	0.065	0.01	55	0.115	0.025
4	0.07	0.015	09	0.110	0.02
5	0.075	0.02	70	0.115	0.02
9	0.075	0.02	80	0.113	0.02
7	0.090	0.015	06	0.110	0.022
8	0.092	0.015			
9	0.095	0.02			
10	0.085	0.05			

Table C-3. Results of Soil Gas Permeability Test at Monitoring Point H1-MPC

	Pressure ("H	("H ₂ O) by Depth		Pressure ("H,O) by Depth	O) by Depth
Time (min)	3.5′	6.0′	Time (min)	3.5′	,0'9
0	0	0	20	0.01	0.02
—	0	0.01	23	0.02	0.015
2	0.01	0.015	26	0.03	0.035
3	0.01	0.03	29	0.03	0.015
4	0.02	0.02	32	0.02	0.02
5	0.03	0.025	35	0.025	0.015
9	0.02	0.03	38	0.015	0.02
7	0.03	0.01	41	0.025	0.15
8	0.01	0.015	44	0.01	0.01
6	0.015	0.005	47	0.005	0.015
10	0.01	0.02	50	0.02	0.04
12	0	0	09	0.01	0.02
14	0.02	0.01	70	0.01	0.015
16	0.015	0.025	80	0.01	0.01
18	0.03	0.02	09	0.015	0.01

APPENDIX D BUILDING 1639 IN SITU RESPIRATION TEST DATA

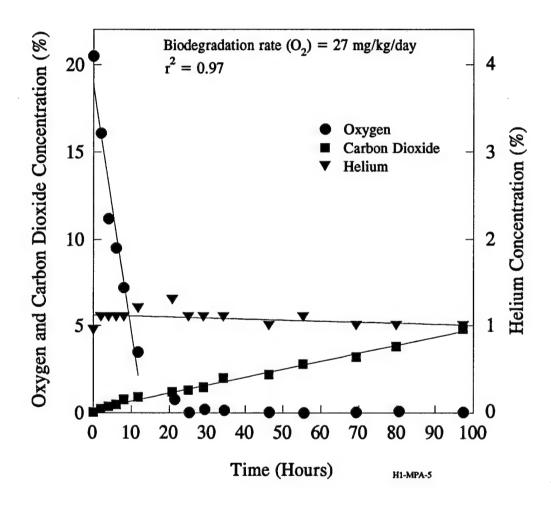


Figure D-1. Oxygen Utilization and Carbon Dioxide Production During the In Situ Respiration Test at Monitoring Point H1-MPA-5.0'

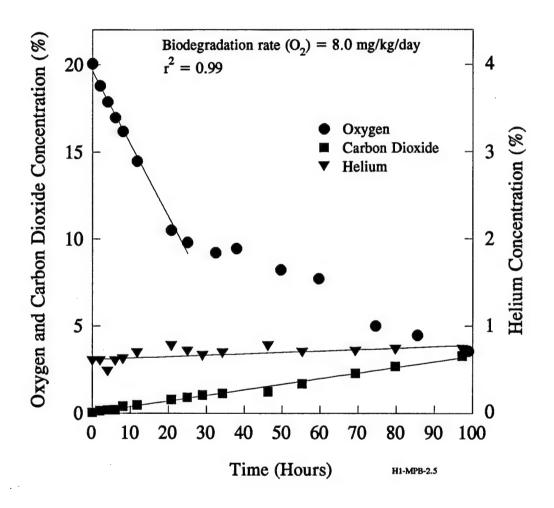


Figure D-2. Oxygen Utilization and Carbon Dioxide Production During the In Situ Respiration Test at Monitoring Point H1-MPB-2.5'

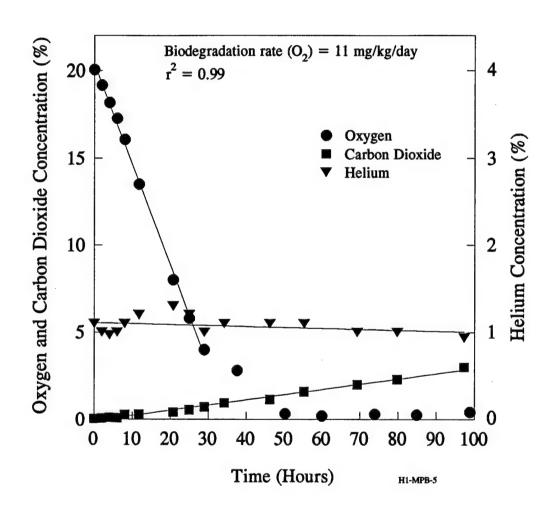


Figure D-3. Oxygen Utilization and Carbon Dioxide Production During the In Situ Respiration Test at Monitoring Point H1-MPB-5.0'

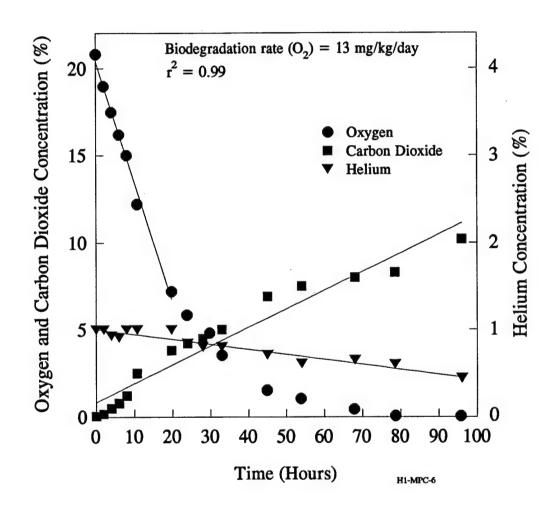


Figure D-4. Oxygen Utilization and Carbon Dioxide Production During the In Situ Respiration Test at Monitoring Point H1-MPC-6.0'

APPENDIX E BUILDING 1812 SOIL GAS PERMEABILITY DATA

Table E-1. Results of Soil Gas Permeability Test at Monitoring Point H2-MPA

	Pressure ("H ₂ O) by Depth	O) by Depth		Pressure ("H ₂ O) by Depth	O) by Depth
Time (min)	2.5′	5.0′	Time (min)	2.5′	5.0′
0	0	0	20	0.55	0
2	0.50	0	25	0.55	0
3	0.54	0	30	0.56	0
4	0.55	0.005	35	0.56	0
5	0.55	0	40	0.56	0
9	0.55	0	90	0.57	0
7	0.55	0	09	85.0	0
8	0.55	0	75	0.58	0
6	0.55	0	06	65.0	0
12	0.55	0			
16	0.55	0.55			
18	0.55	0			

Table E-2. Results of Soil Gas Permeability Test at Monitoring Point H2-MPB¹

	Pressure ("H	ure ("H ₂ O) by Depth		Pressure ("H ₂ O) by Depth	O) by Depth
Time (min)	2.5′	5.0′	Time (min)	2.5′	5.0′
0	0	0	14	0.01	0.01
1	0	0	16	0	0.005
2	0	0	18	0.005	0.005
3	0	0.005	20	0	0
4	0	0	25	0	0
5	0	0	30	0.005	0.005
9	0	0	35	0	0
7	0	0	40	0.005	0.005
8	0	0	50	0.005	0.005
6	0	0	09	0.005	0.005
10	0	0	75	0.005	0.005
12	0.005	0.005			

Pressure readings were not collected from monitoring point H1-MPB-7.0'.

Table E-3. Results of Soil Gas Permeability Test at Monitoring Point H2-MPC

	Press	Pressure ("H ₂ O) by Depth	Depth		Pressi	Pressure ("H ₂ O) by Depth	Depth
Time (min)	2.5′	4.5′	6.0′	Time (min)	2.5′	4.5′	.0.9
1	0	0>	0>	16	0	<0	0>
2	<0>	0>	0	18	0	0>	0>
3	<0>	0>	<0	20	0	<0>	0>
4	<0>	0>	<0	25	0	0>	0>
5	0>	0>	<0	30	0	0	0
9	0	<0>	<0	35	0	0	0
7	0	<0>	<0	40	0	0	0
∞	0	0>	<0	50	0	0	0
6	0	<0>	<0	09	0	0	0
10	0	<0>	<0	70	0	0	0
12	0	0>	<0	80	0	0	0
14	0	<0>	<0>	06	0	0	0